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Method for safety evaluation of chemical compound using recombinant yeast expressing human cytochrome P450.

There is disclosed a method for evaluation of the safety of a chemical compound, which includes the steps of: (a) reacting a chemical compound with recombinant yeast cells expressing, or in other words producing, human cytochrome P450 molecular species P450 1A2, P450 2C9, P450 2E1 and P450 3A4 together with a yeast NADPH-P450 reductase, which may be in the form of a fused enzyme with each of said human cytochrome P450 molecular species, or with the cell free extracts of the yeast cells; and (b) analyzing the resulting metabolite to determine the safety of the compound. According to this method, it can be determined whether a test compound will be converted into a carcinogenic or mutagenic form through the metabolism in the human liver, and whether the test compound or its metabolite has mutagenicity.

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The present invention relates to a method for evaluation of the safety of a chemical compound using recombinant yeasts expressing human cytochrome P450.

The cytochrome P450 is an enzyme catalyzing the mono-oxygenation of a substance in the human liver.

It is known that recombinant human cells expressing heterogeneous human cytochrome P450 species have been used for determination of metabolisms and toxicities of chemical substances. However, this method is unsatisfactory as a method of evaluation of the safety of chemical compounds partly because the kinds of the human cytochrome P450 species expressed by the cells and the levels of the expression are so limited that the amount of metabolite obtained is not enough for determination of the metabolism and toxicity, and partly because it requires not only a high density culture technique but a high cultivation cost. Accordingly, there has been a great demand for developing an advantageous method.

As a result of the extensive study, the present inventors have found that yeasts are particularly suitable as hosts for production of human cytochrome P450 and yeast NADPH-P450 reductase to be used in in vitro determination of metabolisms and toxicities of chemical substances because yeasts grow so rapidly and can stably express both the human cytochrome P450 and yeast NADPH-P450 reductase at high expression levels to provide sufficient amounts of the metabolites in a short period of time, thereby enabling a precise and quick analysis of the metabolites.

Moreover, they have also found that, despite that there are a considerable number of human cytochrome P450 molecular species, the human metabolic system for chemical compounds can be reproduced in vitro when at least four human cytochrome P450 molecular species, i.e., human cytochrome P450 1A2, P450 2C9, P450 2E1 and P450 3A4, are combined.

Thus, the present invention provides a method for evaluation of the safety of a chemical compound, which comprises the steps of:

(a) reacting a chemical compound with recombinant yeast cells expressing, or in other words producing, human cytochrome P450 molecular species P450 1A2, P450 2C9, P450 2E1 and P450 3A4 together with a yeast NADPH-P450 reductase, which may be in the form of a fused enzyme with each of said human cytochrome P450 molecular species, or with the cell free extracts of the yeast cells; and

(b) analyzing the resulting metabolite to determine the safety of the compound.

The present invention further provides a method for determination of the human metabolite of a chemical compound, which comprises the steps of:

- (a) reacting a chemical compound with recombinant yeast cells producing human cytochrome P450 molecular species P450 1A2, P450 2C9, P450 2E1 and P450 3A4 together with a yeast NADPH-P450 reductase, which may be in the form of a fused enzyme with each of said human cytochrome P450 molecular species, or with cell free extracts of the yeast cells; and
- (b) identifying the resulting metabolite.

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- Figs. 1 to 4 show various primers for cloning human P450 genes.
- Fig. 5 shows a synthetic linker for human P450 gene cloning.
- Fig. 6 shows a method of constructing yeast expression plasmids for human P450 1A2.
- Fig. 7 shows a method of constructing yeast expression plasmids for human P450 2C9.
- Fig. 8 shows a method of constructing yeast expression plasmids for human P450 2E1.
- Fig. 9 shows a method of constructing yeast expression plasmids for human P450 3A4.
- Fig. 10 shows a method of constructing yeast expression plasmids for human P450 1A1.
- Fig. 11 shows a method of constructing yeast expression plasmids for human P450 2A6.
- Fig. 12 shows a method of constructing yeast expression plasmids for human P450 2B6.
- Fig. 13 shows a method of constructing yeast expression plasmids for human P450 2C8.
- Fig. 14 shows a method of constructing yeast expression plasmids for human P450 2C18. Fig. 15 shows a method of constructing yeast expression plasmids for human P450 2C19.
- Fig. 16 shows a method of constructing yeast expression plasmids for human P450 2D6.
- Fig. 17 shows a method of constructing a yeast expression plasmid containing an artificial fused enzyme gene.
  - Fig. 18 shows a method of constructing a yeast expression plasmid using a GAPDH promoter.

According to the present invention, it can be determined whether a test compound will be converted into a carcinogenic or mutagenic form through the metabolism in the human liver, and whether the test compound or its metabolite has mutagenicity.

Thus, the present invention provides a method for evaluation of safety of a chemical compound, and a method for determination of the human metabolite of a chemical compound.

### Human Cytochrome P450 and Their Genes

The yeasts capable of expressing, or producing, said enzymes can be obtained by transforming them with expression plasmids containing genes encoding said enzymes with a conventional recombinant DNA method.

The human P450 molecular species to be used in the present invention include at least four human cytochrome P450 molecular species, i.e., human cytochrome P450 1A2, P450 2C9, P450 2E1 and P450 3A4. The genes encoding these essential human cytochrome P450 molecular species and yeast NADPH-P450 reductase are reported in Nucleic Acids Res., 14, 6773-6774, 1986; J. Biochem., 102, 1075-1082, 1987; J. Biol. Chem., 261, 16689-16697, 1986; DNA, 7, 79-86, 1988; and J. Biochem., 103, 1004-1010, 1988.

Although the kinds of P450 molecular species present in human liver vary among the race and individuals, the combination of said human P450 molecular species includes at least about 85% (molar ratio) of the total amount of the human P450 molecular species present in the human liver. Hence, the present method using the said combination of human P450 molecular species can accurately reproduce the human liver metabolism in vitro.

The combination of these P450 molecular species may optionally be varied, taking into account of the amounts of these P450 molecular species in the liver: the amount of P450 3A4 present in the human liver is about 35±10% of the total amount of the human P450 molecular species; P450 2C9 about 25±10%; P450 1A2 about 23±10%; and P450 2E1 about 17±10%.

In addition to the above-mentioned combination, human P450 molecular species P450 2A6, P450 2C19 and/or P450 2D6 (Biochemistry, 29, 1322-1329, 1990; Biochemistry, 30, 3247-3255, 1991; Am. J. Hum. Genet., 45, 889-904, 1989) may also be added. In this case, the combined human P450 molecular species covers at least 90% of the total amount of the human P450 molecular species present in the human liver.

The in vitro human metabolic system that reproduces accurately the human metabolism of a chemical compound, and can represent the differences among races and individuals can be obtained when these human P450 molecular species are properly combined, taking into account of the amounts of these species in the liver.

Furthermore, at least one human cytochrome P450 molecular species selected from the group of P450 1A1, P450 2B6, P450 2C8 and P450 2C18 (Science, 228, 80-83, 1985; Biochemistry, 28, 7340-7348, 1989; Nucleic Acids Res., 15, 10053-10054, 1987; Biochemistry, 30, 3247-3255, 1991) may be added to said human cytochrome P450 molecular species to reproduce in vitro the metabolism of the human liver more accurately.

The nucleotide sequences coding for the human P450 molecular species are disclosed in SEQ ID NOs: 1 to 38.

### Cloning of Genes

The genes coding for the human cytochrome P450 molecular species are known and can be obtained by the conventional cloning methods.

For example, they may be obtained by:

- (i) preparing a mRNA fraction containing the mRNA of the gene coding for human cytochrome P450 molecular species;
- (ii) preparing a cDNA from the mRNA fraction using reverse transcriptase;
- (iii) preparing a cDNA library by inserting said cDNA into a pharge vector or a plasmid vector; and
- (iv) cloning the gene coding for the human cytochrome P450 molecular species from the cDNA library obtained above or from a commercially available human liver-derived cDNA library using a DNA fragment having an identical sequence to some part of the desired gene or an antibody reactive to the protein produced by the gene.

The gene may also be obtained from the above-described cDNA library by the PCR method.

The gene coding for yeast NADPH-P450 reductase may be obtained by the same method as used for cloning of the genes coding for human P450 molecular species. More specifically, the gene may be obtained by such a known method as described in the Japanese Patent Laid-open Publication No. 62-19085.

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### Construction of Yeast Expression Plasmids

The yeasts capable of expressing said enzymes can be obtained by transforming them with expression plasmids containing genes encoding said enzymes with a conventional recombinant DNA method.

The yeast expression plasmid having a gene coding for human P450 molecular species and a gene coding for the yeast NADPH-P450 reductase can be constructed by using a conventional recombinant DNA method.

As to the promoter to be used for construction of the expression plasmids for the yeast of the present invention, there is no particular restriction so long as the promoter can be used in usual expression systems for yeasts, and a promoter of a yeast alcohol dehydrogenase gene (hereinafter referred to as ADH promoter), glyceraldehyde-3-phosphate dehydrogenase promoter (hereinafter referred to as GAPDH promoter), and phosphoglycerate kinase (hereinafter referred to as PGK promoter) are preferably used in the present invention.

The ADH promoter can be prepared by a usual genetic engineering method, for example, from a yeast expression vector pAAH5 possessing a yeast ADH1 promoter and terminator ("Methods in Enzymology" by Ammerer et al., vol.101, pp.192-201). The yeast ADH1 promoter is described in the U.S. Patent No. 299,733 to Washington Research Foundation and it requires patent license from the patentee in a case of using the same for an industrial or commercial purpose.

The yeast expression plasmid having both a gene coding for human P450 molecular species and a gene coding for the yeast NADPH-P450 reductase can be constructed by, for example, inserting an Notl tragment prepared from yeast expression vector pAAH5N possessing the ADH promoter and terminator (Japanese Patent Laid-open Publication No. 2-211880) to an Notl site of plasmid pARRN possessing a gene coding for yeast NADPH-P450 reductase (Japanese Patent Laid-open Publication No. 2-211880) and then inserting cDNA coding for the human P450 molecular species to the HindIII site of the thus obtained plasmid pAHRR. Moreover, a vector obtained by exchanging a Hind III site of pAAH5N with other restriction enzyme site may be used for the same purpose.

In the present invention a gene coding for an artificial fused enzyme comprising human cytochrome P450 molecular species and yeast NADPH-P450 reductase can also be used. The artificial fused enzyme can catalyze mono-oxygenation reaction and the efficiency of the electron transfer from NADPH is so improved that the activity of the mono-oxygenation reaction is much enhanced. Accordingly, a great amount of metabolic products can be obtained in a shorter period of time, enabling accurate analysis.

The fused gene comprises a gene coding for the human cytochrome P450 molecule on the 5'-terminal and a gene coding for the yeast NADPH-P450 reductase on 3'-terminal.

The gene coding for such an artificial fused enzyme can be constructed by ligating a gene coding for a human cytochrome P450 species and a gene coding for yeast NADPH-P450 reductase by a conventional recombinant DNA method, and the constructed gene is usually inserted to the Hind III site of the yeast expression vector pAAH5N having ADH promoter and ADH terminator described in the Japanese Patent Laid-open Publication No. 2-211880.

#### Transformation of Yeast

The yeast cells expressing the human P450 molecular species and yeast NADPH-P450 reductase or yeast cells expressing an artificial fused enzyme comprising human P450 molecular species and NADPH-P450 reductase can be obtained by introducing the thus constructed yeast expression plasmid into a yeast by a known method such as a protoplast method or a method using alkaline metal salt (LiCI).

In the present invention, two or more expression plasmids may optionally be introduced into a single strain of yeast so that the yeast can express two or more molecular species simultaneously.

As the hosts, Saccharomyces cerevisiae is used in the method of the present invention, in particular, Saccharomyces cerevisiae AH22 (ATCC 38626) is preferably used.

### Reaction of Test Compound

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In the method of the present invention, a test compound is reacted with a mixture of at least said four human P450 molecular species, or separately with each of the said four human P450 molecular species in the presence of the yeast NADPH-P450 reductase.

Alternatively, it may be first reacted with one or more of the essential human P450 molecular species, and then with a mixture of, or separately with the rest of them; each of the reactions is carried out in the presence of the yeast NADPH-P450 reductase.

The reaction is carried out by reacting a test compound with the yeast obtained by the transformation with an expression plasmid containing a gene encoding a human P450 molecular species and a gene encoding yeast NADPH-P450 reductase, or a fused gene encoding a fused enzyme of a human P450 molecular species and a yeast NADPH-P450 reductase, or with the cell free extracts of the yeast cells.

In the reaction of a test compound with the enzymes of the present invention, living yeast cells and their cell free extracts are usually used.

As the cell free extracts, subcellular fraction of cells containing microsomal fractions, or fractions containing both microsome and cytoplasm is used. The cell free extracts or fractions can be prepared, for example, by a known method (DNA, Vol.4, No. 3, pp.203-210 (1985)).

However, the present invention can be preferably carried out with the cell free extracts, especially with microsomal fractions of the cells. But, when biological analytic method is used to determination of the mutagenicity or carcinogenicity, fractions containing microsome and cytoplasm are preferably used.

The reaction can be conducted by adding a test compound to a culture solution or a buffer solution of yeast cells or cell free extracts, and the resultant solution is usually incubated at a temperature, for example, at about 10 °C to 40 °C, for about 0.1 to 48 hours.

The amounts of the yeast cells or cell free extracts and the compound vary depending on the conditions such as reaction temperature, reaction time and the kind of the test compound to be used.

For instance, the amount of the yeast cells to be used in the solution is preferably from about  $10^5$  to about  $10^{10}$  per 1 ml of the solution, preferably, from about  $10^7$  to about  $10^8$  per 1 ml of the solution. When cell free extracts are used, from about  $10^{10}$  to about  $10^{15}$  of P450 molecules per 1 ml of the solution, preferably from about  $10^{12}$  to about  $10^{13}$  of P450 molecules per 1 ml of the solution is usually used.

The amount of the compound to be added is preferably within a range of from about 0.01 µmol to about 1µmol per 1 ml of the solution.

The above ranges may be optionally varied, if necessary.

### **Determination of Metabolites**

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The metabolites present in the reaction solution can then be subjected to elucidation of the chemical structures and the measurement of their amounts. The analysis of the chemical structure can be conducted by known methods ("Guide to Apparatus Analysis (2)", edited by Jiro Shiokawa et al., (revised edition) first print, issued from Kagaku Dojin (1985); "Spectral Identification for Organic Compound" by R.M. Silverstein, fourth edition, third print, issued from Tokyo Kagaku Dojin (1984)).

From the results of the analysis of the metabolites, it can be determined whether the tested compound will be detoxicated or metabolized into a carcinogen in the human liver when administered.

### Determination of Toxic Effects of Metabolites

The toxic effects, in particular mutagenicity, of the resulting metabolites can be determined by a conventional biological analytic method such as the Ames Test. For example, the metabolites present in the reaction solution are allowed to react with mutant bacteria such as histidine requiring Salmonella strain (Salmonella typhimurium (his-)), or tryptophan requiring Escherichia coil (Escherichia coil (trp-)), and then determine whether the metabolites cause the back mutation of the bacteria whether the colonies of revertant which is not requiring the amino acid (His+ or Trp+) are formed, and, if formed, how many colonies. In place of the bacteria, mammalian cells such as MCL-5 cells, which are sensitive to cell toxicity of a chemical compound (U.S. Patent No. 4,532,204), can be used.

In this method, the compounds that cause the back mutation will be judged to be mutagenicity test-positive.

It is also possible to simultaneously proceed the step (a) of reacting the test compound with the yeast cells or the cell free extracts, and the step (b) of analyzing the metabolites present in the reaction solution.

The mutagenicity of arylamine derivatives, which are known to be metabolized by the liver into a mutagens, can be examined by the biological analytic method. For example, the mutagenicity of 2-aminoanthrathene can be detected at the concentration of about 0.1  $\mu$ g of 2-aminoanthrathene when 20 pmol of P450 1A2, which is active specifically to 2-aminoanthrathene, is used (Table 1).

In the present invention, a metabolic probe for a human P450 molecular species can be obtained.

If a certain chemical compound is converted by a particular human P450 molecular species into a specific metabolite, the amount of such a human P450 molecular species can be determined by detecting such a metabolite in excretions such as blood or urine of a living body who has been administered the compound, and such a compound is called a metabolic probe.

In the present invention, such a metabolic probe can be obtained by screening the metabolites obtained by reacting chemical compounds with the yeasts of the present invention.

### Examples

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The present invention will be further illustrated by the following examples, which are not to be construed to limit the scope thereof.

## Preparation of cDNA coding for human P450 molecular species

cDNA coding for human P450 molecular species were obtained from commercially available human liver cDNA library (Clontech Co.) by the PCR method using primers for cloning human P450 genes as shown in Figs. 1 to 4, and a method using a synthetic linker for human P450 gene cloning as shown in Fig. 5. Thus obtained nucleotide sequences for the cDNA and the deduced amino acid sequences are shown in the sequence listing.

Relationship between SEQ ID NOs and human P450 molecular species are as follows:

1. The essential human cytochrome P450 molecular species for the present invention.

(1) SEQ ID NO: 1	1A2
(2) SEQ ID NO: 3	2C9
(3) SEQ ID NO: 5	2E1
(4) SEQ ID NO: 7	3A4

#### 2. Auxiliary Human cytochrome P450 molecular species

	(1) SEQ ID NOs: 9, 11 and 13	1A1
	(2) SEQ ID NOs: 15 and 17	2A6
	(3) SEQ ID NO: 19	2B6
	(4) SEQ ID NOs: 21, 23 and 25	2C8
Ì	(5) SEQ ID NO: 27	2C18
ĺ	(6) SEQ ID NO: 29	2C19
	(7) SEQ ID NOs: 31, 33, 35 and 37	2D6

## Construction of yeast expression plasmids: p1A2 and p1A2R

Fig. 6 shows a method of constructing yeast expression plasmids for human P450 1A2. The protein coding region of P450 1A2 gene of about 1.5 kb excluding about 40 bp at the 5'-terminal was amplified by the PCR method using the primers shown in Fig. 1. The resultant fragment of about 1.5 kb was cleaved with SacI and sub-cloned to a pUC118 vector. About 40 bp at the 5'-terminal was chemically synthesized as the linkers shown in Fig. 5 and sub-cloned between the HindIII and SacI sites of the pUC118 vector. The plasmid having the 1.5 kb fragment was digested by HindIII, blunted, and then ligated with an EcoRI linker. The EcoRI-SacI fragment was prepared from the resulting plasmid and ligated into the plasmid containing the 5'-terminal 40 bp. Then, it was treated with EcoRI and blunted. A HindIII linker was inserted into the blunted fragment. The obtained fragment then cleaved with HindIII was inserted into pAAH5N and pAHRR to construct a yeast expression plasmid p1A2 for human P450 1A2, and a yeast expression plasmid p1A2R for simultaneous expression of human P450 1A2 and yeast NADPH-P450 reductase.

## Construction of yeast expression plasmids: p2C9 and p2C9R

Fig. 7 shows a method of constructing yeast expression plasmids for human P450 2C9. The protein coding region of 450 2C9 gene was divided into two fragments of about 0.9 kb and about 0.6 kb, and the fragments were amplified by the PCR method using the primers shown in Fig. 1. The resultant fragment of about 0.9 kb was cleaved with Pstl and sub-cloned to a pUC B vector, which was prepared by exchanging the cloning site located between the two Hind III sites, one of which was obtained by converting the EcoRI site of pUC19, with the following cloning sites:

EcoF	RI	SpeI	PstI	Bam	HI	KpnI	Hind	III
HindIII	XbaI	SphI	:	SalI	SmaI		SacI	
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The fragment of about 0.6 kb was incorporated between the Xbal and Pstl sites of the plasmid having the 0.9 kb fragment to ligate the two segments. The Kpnl site of the plasmid was blunted. An Xbal linker was inserted to the blunted plasmid. The Xbal fragment containing the coding region was cut out from the resultant fragment. A modified pUC vector, pUCAN, was constructed by replacing the EcoRl and HindIll sites with Notl sites, followed by insertion of the Notl fragment prepared from pAAH5N between the two Notl sites. The HindIll site of pUCAN vector having the ADH promoter and terminator regions in the pUC vector was blunted and inserted into pUCANX introduced with the Xbal linker. The obtained plasmid was cleaved with Notl and inserted into pAAH5N and pAHRR treated in a similar manner with Notl to construct a yeast expression plasmid p2C9 for human P450 2C9, and a yeast expression plasmid p2C9R for simultaneous expression of human P450 2C9 and yeast NADPH-P450 reductase.

### Construction of yeast expression plasmids: p2E1 and p2E1R

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Fig. 8 shows a method of constructing yeast expression plasmids for human P450 2E1. The protein coding region of P450 2E1 gene was divided into two fragments of about 0.5 kb and about 1.0 kb, both of which were amplified by the PCR method using the primers shown in Fig. 1. The resultant fragment of about 0.5 kb was cleaved with EcoRI and BamHI and sub-cloned to a pUC118 vector. Then the fragment of about 1.0 kb was incorporated between the BamHI and SphI sites to ligate the two fragments. This was cleaved with EcoRI, and SphI, and inserted into pUC B first and then cut out with HindIII. The resultant fragment was inserted into pAAH5N and pAHRR vectors to construct a yeast expression plasmid p2E1 for human P450 2E1, and a yeast expression plasmid p2E1R for simultaneous expression of human P450 2E1 and yeast NADPH-P450 reductase.

## Construction of yeast expression plasmids: p3A4 and p3A4R

Fig. 9 shows a method of constructing yeast expression plasmids for human P450 3A4. The protein coding region of P450 3A4 gene was divided into two fragments of about 0.6 kb and about 0.9 kb, both of which were amplified by the PCR method using the primers shown in Fig. 2. The resultant fragment of about 0.6 kb was cleaved with SacI and sub-cloned to a puC118 vector. Subsequently, it was cleaved with EcoRI and blunted. An XbaI linker was ligated to the blunted fragment. The fragment of 0.9 kb was cleaved with XbaI and SacI, and incorporated to the resultant fragment above, thus the two fragments were ligated. After cleaving the plasmid with SphI, it was blunted. An XbaI linker was ligated to the blunted fragment, from which the XbaI segment was cut out and inserted to an XbaI site of pUCANX. This was cut out with NotI and inserted into pAAH5N and pAHRR treated in a similar manner with NotI. Thus a yeast expression plasmid p3A4 for human P450 3A4, and a yeast expression plasmid p3A4R for simultaneous expression of human P450 3A4 and yeast NADPH-P450 reductase were constructed.

## Construction of yeast expression plasmids: p1A1 and p1A1R

Fig. 10 shows a method of constructing yeast expression plasmids for human P450 1A1. The coding region for P450 1A1 protein was divided into two fragments of about 1.0 kb and about 0.5 kb and the resultant fragments were amplified by the PCR method using the primers shown in Fig. 2. Thus obtained fragment of about 1.0 kb was cleaved with Xbal and Sacl and sub-cloned to a PUCA vector, which was prepared by exchanging the cloning site located between the two HindIII sites, one of which was obtained by converting the EcoRI site of pUC19, with the following cloning sites:

	Хb	aI	SpeI	Pst	I	BamHI	KpnI	Hino	III
	HindIII	EcoRI	S	phI	SalI	Sma I	- -	SacI	
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The amplified fragment of about 0.5 kb was sub-cloned into the HinclI site of a pUC 19 vector and the resultant plasmid was then cleaved with Sacl. The cleaved fragment was ligated with the plasmid having the 1.0 kb fragment. After cutting out the coding region from the thus obtained 1A1 gene with HindIII, the fragment was inserted to the HindIII site of the yeast expression vector pAH5N having ADH promoter and terminator regions, and to the same site of vector pAHRR for simultaneous expression of P450 and yeast NADPH-P450 reductase of which gene is located upstream of the P450 gene. Thus yeast expression plasmid p1A1 for human P450 1A1 and yeast expression plasmid p1A1R for simultaneous expression of human P450 1A1 and yeast NADPH-P450 reductase were constructed.

In addition two kinds of human P450 1A1 gene fragments which were different only in a small portion of the nucleotide sequence were obtained in a similar manner and used to construct two kinds of yeast expression plasmid for human P450 1A1, p1A1 Variant 1 and p1A1 Variant 2, and two kinds of plasmids for simultaneous expression of human P450 1A1 and yeast NADPH-P450 reductase, p1A1R Variant 1 and p1A1R Variant 2.

### 15 Construction of yeast expression plasmids: p2A6 and p2A6R

Fig. 11 shows a method of constructing yeast expression plasmids for human P450 2A6. A protein coding region of P450 2A6 gene was divided into two fragments of about 0.6 kb and about 0.9 kb, both of which were amplified by the PCR method using the primers shown in Fig. 2 to yield two kinds of human P450 2A6 gene fragments which were different only in a small portion of the nucleotide sequence. The resultant fragment of about 0.6 kb was cleaved with Xbal and Hincll, and sub-cloned to a pUC A vector. Then the fragment of 0.9 kb was incorporated between the Hincll and Kpnl sites to ligate the two fragments. The obtained fragment was cleaved with Hindlll and inserted into pAAH5N and pAHRR to construct two kinds of yeast expression plasmid for human P450 2A6, p2A6 and p2A6 Variant 1, and two kinds of yeast expression plasmid for simultaneous expression of human P450 2A6 and yeast NADPH-P450 reductase, p2A6R and p2A6R Variant 1.

## Construction of yeast expression plasmids: p2B6 and p2B6R

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Fig. 12 shows a method of constructing yeast expression plasmids for human P450 2B6. The entire protein coding region of P450 2B6 gene was amplified by the PCR method using the primers shown in Fig. 3. The resultant fragment was cleaved with Xbal and BamHI and sub-cloned to pUC A. The resulting plasmid was partially digested with HindIII, and inserted into pAAH5N and pAHRR vectors to construct a yeast expression plasmid p2B6 for human P450 2B6, and a yeast expression plasmid p2B6R for simultaneous expression of human P450 2B6 and yeast NADPH-P450 reductase.

### Construction of yeast expression plasmids: p2C8 and p2C8R

Fig. 13 shows a method of constructing yeast expressed plasmids for human P450 2C8. The entire protein coding region of the P450 2C8 gene was amplified by the PCR method using the primers shown in Fig. 3 to yield three kinds of P450 2C8 genes which were different only in a small portion of the nucleotide sequence. The resultant fragments were partially digested with Xbal, and sub-cloned to pUC A. The fragment was cleaved with HindIII and inserted into pAAH5N and pAHRR vectors to construct three kinds of yeast expression plasmids p2C8, p2C8 Variant 1 and p2C8 Variant 2 for human P450 2C8, and three kinds of yeast expression plasmids, p2C8R, p2C8R Variant 1 and p2C8R Variant 2 for simultaneous expression of human P450 2C8 and yeast NADPH-P450 reductase.

## Construction of yeast expression plasmids: p2C18 and p2C18R

Fig. 14 shows a method of constructing yeast expression plasmids for human P450 2C18. The protein coding region of P450 2C18 gene was divided into two segment of about 1.4 kb and about 0.9 kb, then the both fragments were amplified by the PCR method using the primers shown in Fig. 3. The amplified fragment of about 1.4 kb was cleaved with Pstl and sub-cloned to a pUC A vector. The fragment of about 0.9 kb was incorporated between the Xbal and Pstl sites to ligate the two fragments. After cleaving the plasmid with Smal, an Xbal linker was introduced. Then an Xbal fragment was prepared and inserted into the Xbal site of pUCANX. It was cleaved with Notl and inserted into pAAH5N and pAHRR treated in a similar manner with Notl to construct a yeast expression plasmid p2C18 for human P450 2C18, and a yeast expression plasmid p2C18R for simultaneous expression of human P450 2C18 yeast and NADPH-P450

reductase.

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## Construction of yeast expression plasmids: p2C19 and p2C19R

Fig. 15 shows a method of constructing yeast expression plasmids for human P450 2C19. Fragments a, b and c for the protein coding region of P450 2C19 gene were amplified by the PCR method using the primers No. 1, No. 2, No. 3 and No. 4, No.5 and No. 6, and No.5 and No.7 defined by SEQ ID NOs: 39-45, respectively.

Fragments e and f for the protein coding region of human cytochrome P450 2C19 were also amplified against human cytochrome P450 2C9 gene by the PCR method using the primers No. 8 to 21 having nucleotide sequences with some mutations shown by SEQ ID NOs: 46 to 59. A fragment d for the linker Nos. 1 and 2 having nucleotide sequences shown by SEQ ID NOs: 60 and 61 was obtained by directly synthesizing the DNA to cover the rest of the protein coding region of the human P450 2C19 gene. Thus the fragments covering the whole protein coding region of the human cytochrome P450 2C19 were obtained.

After the fragments a and b were treated with Xhol and BamHl, and with BamHl and Pstl, both fragments were simultaneously inserted between the Xhol and Pstl sites of the Blue Script(+). The fragment e was treated with Xbal and Xhol and inserted to the Xbal and Xhol sites of the plasmid having the fragments a and b to give a plasmid having the fragments a, b and e.

After the fragment c was treated with Pstl and Kpnl, the resulting fragment was simultaneously inserted with the linker fragment d between the Pstl and EcoRl sites of the Blue Script(+). The resultant plasmid was cut with Pstl and EcoRl to give a fragment containing the fragments c and d. Then this fragment was simultaneously inserted between the fragment f treated with EcoRl to the Pstl and Hincll sites of the aforementioned plasmid containing the fragment a, b and e. Thus a plasmid having the whole coding region of the human cytochrome P450 2C19 gene was constructed. The constructed plasmid was cut with Hindlll and the resultant fragment was inserted to pAAH5N and pAHRR both of which were treated with Hindlll to give a yeast expression plasmid p2C19 for expressing the human P450 2C19 and a yeast expression plasmid p2C19R for simultaneous expression of the human P450 2C19 and yeast NADPH-P450 reductase.

o SEQ ID NOs and primer Nos. are as follows:

SEQ ID No: 39	Primer No. 1
SEQ ID NO: 40	Primer No. 2
SEQ ID NO: 41	Primer No. 5
SEQ ID NO: 42	Primer No. 4
SEQ ID NO: 43	Primer No. 5
SEQ ID NO: 44	Primer No. 6
SEQ ID NO: 45	Primer No. 7
SEQ ID NO: 46	Primer No. 8
SEQ ID NO: 47	Primer No. 9
SEQ ID NO: 48	Primer No. 10
SEQ ID NO: 49	Primer No. 11
SEQ ID NO: 50	Primer No. 12
SEQ ID NO: 51	Primer No. 13
SEQ ID NO: 52	Primer No. 14
SEQ ID NO: 53	Primer No. 15
SEQ ID NO: 54	Primer No. 16
SEQ ID NO: 55	Primer No. 17
SEQ ID NO: 56	Primer No. 18
SEQ ID NO: 57	Primer No. 19
SEQ ID NO: 58	Primer No. 20
SEQ ID NO: 59	Primer No. 21
SEQ ID NO: 60	Linker No. 1
SEQ ID NO: 61	Linker No. 2
	SEQ ID NO: 40 SEQ ID NO: 41 SEQ ID NO: 42 SEQ ID NO: 43 SEQ ID NO: 44 SEQ ID NO: 45 SEQ ID NO: 45 SEQ ID NO: 46 SEQ ID NO: 47 SEQ ID NO: 48 SEQ ID NO: 50 SEQ ID NO: 51 SEQ ID NO: 52 SEQ ID NO: 53 SEQ ID NO: 55 SEQ ID NO: 55 SEQ ID NO: 55 SEQ ID NO: 55 SEQ ID NO: 57 SEQ ID NO: 58 SEQ ID NO: 59 SEQ ID NO: 59 SEQ ID NO: 60

### Construction of yeast expression plasmids: p2D6 and p2D6R

Fig. 16 shows a method of constructing yeast expression plasmids for human P450 2D6. The protein coding region of 1.3 kb excluding about 200 bp at the 5'-terminal of P450 2D6 gene was divided into two fragments of about 0.4 kb and about 0.9 kb, and the both fragments were amplified by the PCR method. The resultant fragment of about 0.9 kb was cleaved with Kpnl and sub-cloned to pUC A. For the 200 bp on the 5'-terminal, three synthetic linkers shown in Fig. 5 were used and two linkers on the 5'-terminal were incorporated into Xbal and Pstl sites of a Blue Script(+) vector and then other linkers were incorporated into Smal and Pstl sites. Then fragment of about 0.4 kb obtained by the PCR method was incorporated into the Pstl and Hincll sites of the plasmid and then cleaved with NspV and Xbal. The resultant fragment was inserted into the plasmid containing the 0.9 kb fragment to ligate the coding region. This was cleaved with HindIII and inserted into pAAH5N and pAHRR vectors to construct a yeast expression plasmid p2D6 for human P450 2D6, and a yeast expression plasmid p2D6R for simultaneous expression of human P450 2D6 and yeast NADPH-P450 reductase.

Then three kinds of human P450 2D6 gene fragments which were different only in a small portion of the nucleotide sequence were obtained in a similar manner as described above and used to construct two kinds of yeast expression plasmids for human P450 2D6, p2D6 Variant 1, p2D6 Variant 2 and p2D6 Variant 3, and three kinds of yeast expression plasmid 2D6R for simultaneous expression of human P450 2D6 yeast and NADPH-P450 reductase, p2D6R Variant 1, p2D6R Variant 2 and p2D6R Variant 3.

## Construction of yeast expression plasmid containing artificial fused enzyme gene

An expression plasmid was constructed in accordance with Fig. 17. The Xbal-Xhol fragment was amplified with plasmid p3A4 by using the primers shown in Fig. 4. On the other hand, the Xhol-HindIll fragment of about 2.1 kb was obtained from the plasmid pBFCRI (Japanese Patent Application No. 4-209226) and inserted between the Xhol and HindIll sites of a commercial vector Blue Script(+), followed by digestion with restriction enzymes Xhol and Xbal. These two fragments were simultaneously inserted to the Xbal site of the vector pUCAN, which was then digested with Notl to give a fragment of about 5.6 kb. The desired yeast expression plasmid pF3A4 was obtained by ligating the fragment with the Notl fragment of about 10.5 kb obtained from vector pAAH5N (Japanese Patent Laid-open Publication No. 2-211880). The artificial fused enzyme consists of 1156 amino acid residues of which sequence structure comprising, successively, from the N-terminal end, an entire amino acid sequence (503 residues) of human liver cytochrome P450 3A4, a linker-derived sequence (Ala-Arg-Ala), and a sequence of from the 42nd residue to C-terminal of yeast NADPH-cytochrome P450 reductase.

## Preparation of transformed yeast cell

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Saccharomyces cerevisiae AH 22 was inoculated to 1.0 ml of YPD culture medium (1% yeast extract, 2% polypeptone, 2% glucose). After shaken at 30 °C for 18 hours, the yeast cells were collected by centrifugation (5000 x g, 10 min). The resultant cells were suspended in 10 ml of 0.2 M LiCl solution and then centrifuged again (5000 x g, 10 min) to obtain pellets. Then 20  $\mu$ l of 1 M LiCl solution, 30  $\mu$ l of 70% polyethylene glycol 4000 and each 10  $\mu$ l solution containing about 1.0  $\mu$ g of various kinds of yeast expression plasmids for the human P450 molecular species and yeast NADPH-reductase constructed as above were added to the resultant pellets. After sufficiently mixing them, they were incubated at 30 °C for one hour and further stirred after the addition of 140  $\mu$ l of sterilized water. The solution was plated on SD synthetic culture medium (2.0% glucose, 0.67% nitrogen base w/o amino acids, manufactured by Difco Co., 20  $\mu$ g/ml of histidine, 2.0% agar) and incubated at 30 °c for three days. Then transformed yeast cells possessing the yeast expression plasmid described above were selected. In this way, various kinds of yeast cells expressing the human P450 molecular species were prepared.

### Quantitative measurement of human P450 expressed in yeast

Each 200 ml of culture broth of each kind of yeast cells expressing human P450 molecular species and yeast NADPH-reductase simultaneously or expressing an artificial fused enzyme comprising human P450 molecular species and yeast NADPH-reductase prepared as above (SD synthetic culture medium, cell concentration: about 1.5 x 10<sup>7</sup> cells/ml) was used to collect the cells. The collected cells were then suspended in 10 ml of 100 mM potassium phosphate buffer solution (pH 7.0) and centrifuged (5000 x g, 10 min) to obtain pellets. Thus obtained pellets were resuspended in 2.0 ml of 100 mM potassium phosphate

buffer solution (pH 7.0) and 1 ml of each of the solutions were poured into two cuvettes. After bubbling carbon monoxide to a sample cuvette, 5 to 10 mg of dithionite was added to both of the cuvettes, and stirred and then difference spectrum at 400-500 nm was measured to calculate the concentration of P450 present in the yeast. The amount of each kind of human P450 species or an artificial fused enzyme in each kind of transformed yeast cells was at a level from about 10<sup>5</sup> to about 10<sup>6</sup> molecules/cell.

#### Preparation of yeast S-9 Mix fraction, cytoplasmic fraction and microsomal fraction

First, 3.8 liter of each kind of culture broth (SD synthetic culture medium, cell concentration: about 1.0 x 10<sup>8</sup> cells/ml) of yeast cells expressing human P450 molecular species and yeast NADPH-reductase simultaneously or an artificial fused enzyme comprising human P450 molecular species and yeast NADPH-reductase prepared as above was collected and the resultant cells were suspended in 400 ml of a buffer solution A (10 mM Tris-HCl (pH 7.5), 2 M sorbitol, 0.1 mM DTT, 0.2 mM EDTA), to which 160 mg of Zymolyase 100,000 (Zymolyase 100T) was added, and the obtained solution was incubated at 30 °C for 60 min. Spheroplast obtained by centrifugation (5000 x g, 10 min) was suspended in 100 ml of the buffer solution A and then centrifuged (5000 x g, 10 min). Washing the spheroplast by repeating the same centrifugal operation once again, the spheroplast was finally suspended in 200 ml of a buffer solution (10 mM Tris-HCl (pH 7.5), 0.65 M sorbitol, 0.1 mM DTT), which was then subjected to ultrasonic pulverization (50 W, for 5 min). The cell free extracts were centrifuged (9000 x g, 20 min) and supernatants were recovered to obtain a yeast S-9 Mix fraction. Further, the fraction was centrifuged (125,000 x g, 70 min) to collect precipitates which were suspended again into 10 ml of 0.1 M potassium phosphate buffer solution (pH 7.4) to obtain a microsomal fraction. On the other hand, a cytoplasmic fraction was obtained by recovering the supernatants.

#### 25 Construction of yeast expression plasmid using GAPDH promoter and its expression in yeast

Fig. 18 shows a method of constructing a yeast expression plasmid using a GAPDH promoter. A HindIII fragment (about 3.0 kb) obtained from pARRN (described in the Japanese Patent Laid-open Publication No. 2-211880) was inserted into a HindIII site of plasmid pUN, which was obtained by cleaving pUC19 with EcolRI, blunt-ending and ligation with an Notl linker to give pUR. On the other hand, after blunting an Xhol site of plasmid pAAH5 and inserting an Xbal linker, it was cleaved with restriction enzymes Xbal and Sall and the resultant fragment (about 2.2 kb) was inserted to Xbal and Sall sites of pUC19. The three fragments, namely, a fragment (about 2.2 kb) obtained by cleaving the resultant plasmid with Xbal and Pstl, the Xbal-Pstl fragment (about 1.3 kb) cut out from 2 µm DNA of Saccharomyces cerevisiae AH22, and a fragment obtained by cleaving pUR with Pstl were ligated to give a plasmid pURL. Further, the pURL was cleaved with HindIII, blunted and ligated to remove the HindIII site. Then, an Notl fragment (about 1.6 kb) containing GAPDH promoter and terminator (obtained by the method as described in Agric. Biol. Chem., 51, 1641-1647 (1987) and J. Biol. Chem., 267, 16497-16502 (1992)) was ligated to the Notl site of pURL to give a plasmid pURLG. Human P450 2D6 cDNA obtained by the method used for the construction of p2D6 was inserted to a HindIII site of pURLG to obtain a yeast expression plasmid pG2D6R for simultaneous expression of human P450 2D6 and yeast NADPH-P450 reductase. When the plasmid was introduced by the method used in the preparation of transformed yeast cells as above to Saccharomyces cerevisiae AH22, production of human P450 2D6 was observed.

### Metabolism of 7-ethoxycoumarin using transformed yeast cells

7-Ethoxycoumarin was added to each 2 ml of the culture media of the transformed yeast cells expressing (i) human cytochrome P450 molecular species and yeast NADPH-P450 reductase; or (ii) an artificial fused enzyme comprising human cytochrome P450 molecular species and yeast NADPH-P450 reductase (SD synthetic culture medium, cell concentration: about  $2.0 \times 10^7$  cells/ml) so that the final concentration of 7-ethoxycoumarin was 0.5 mM. After incubation at  $30^{\circ}$ C for 2 or 5 hours, supernatants were obtained by centrifugation (5000 x g, 10 min). To the supernatants 62.5  $\mu$ l of 15% TCA (trichloroacetic acid) and 2 ml of chloroform were added and, after well stirring, a chloroform layer was recovered by centrifugation (5000 x g, 10 min), to which 4 ml of 0.01 N NaOH containing 0.1 M NaCl was added and stirred sufficiently and then centrifuged (5000 x g, 10 min). After recovering the supernatants, fluorescence was measured for the supernatant fraction (ex. 366 nm, em 452 nm) to quantitatively measure the reaction product 7-hydroxycoumarin. As a result, 0-deethylation activity for 7-ethoxycoumarin can be observed for all of 11 kinds of the yeast cells expressing the human P450 molecular species. P450 1A1 and P450 2B6

showed strong activity; and P450 1A2, P450 2E1, P450 2A6 and P450 2D6 showed good activity, while P450 2C8, P450 2C9, P450 3A4, P450 2C18 and P450 2C19 showed moderate activity.

### Metabolism of tolbutamide using transformed yeast cells

In the same manner as above, tolbutamide was added to each of the culture solutions of the transformed yeast cells expressing (i) human cytochrome P450 molecular species and yeast NADPH-P450 reductase; or (ii) an artificial fused enzyme comprising human cytochrome P450 molecular species and yeast NADPH-P450 reductase so that the concentration of the compound was 1.0 mM. After incubation at 30 °C for 15 hours, the culture supernatant was then obtained by centrifugation (5000 x g, 10 min). To the supernatant, 2 ml of dichloromethane was added. After sufficient stirring, the dichloromethane layer was recovered by centrifugation (5000 x g, 10 min), and the solvent was evaporated under reduced pressure. The resultant residue was dissolved in 100 µl of acetonitrile, and the solution was analyzed by HPLC under the following conditions. As a result, hydroxylated tolbutamide was detected in the solution of yeast cells expressing human P450 2C8, P450 2C9, P450 2C18 and P450 2C19. The human P450 2C9 showed high activity and 2C19 showed good activity. On the other hand, hydroxylated tolbutamide was not detected in the solution of yeast cells expressing other human P450 than described above.

Conditions for HPLC

Column:

5

20

μBondapak C18 (manufactured by Waters Co.)

Carrier:

10-70% acetonitrile-distilled water (linear concentration gradient for 20 min)

Temperature: Detection:

50 ° C UV 230 nm

Injection amount:

50 μΙ

#### 25 Metabolism of testosterone using transformed yeast cells

In the same manner as above, testosterone was added to each of the culture solutions of the transformed yeast cells expressing (i) human cytochrome P450 molecular species and yeast NADPH-P450 reductase; or (ii) an artificial fused enzyme comprising human cytochrome P450 molecular species and yeast NADPH-P450 reductase so that the concentration of the compound was 0.05 mM. After incubation at 30 °C for 15 hours, the supernatant was obtained by centrifugation (5000 x g, 10 min). Then 2 ml of dichloromethane was added. After sufficient stirring, the solution was centrifuged again (5000 x g, 10 min). The dichloromethane layer was recovered from the separated layer and the solvent was evaporated under reduced pressure. The resultant residue was dissolved in 100  $\mu$ l of acetonitrile, and the solution was analyzed by HPLC under the following conditions. As a result, hydroxylated testosterone was detected for yeast cells expressing human P450 1A1, P450 2C8 and P450 3A4. On the other hand, hydroxylate testosterone was not detected for yeast cells expressing other human P450 than described above.

Conditions for HPLC

Column:

μBondapak C18 (manufactured by Waters Co.)

Carrier:

40

55

20-70% acetonitrile-distilled water (linear concentration gradient for 25 min)

Temperature:

50 · C

Detection:

UV 254 nm

Injection amount:

50 µl

### 45 Metabolism of chlorzoxazone using transformed yeast cells and microsomal fractions thereof

Chlorzoxazone was added to each of the culture solutions of the transformed yeast cells expressing (i) human cytochrome P450 molecular species and yeast NADPH-P450 reductase; or (ii) an artificial tused enzyme comprising human cytochrome P450 molecular species and yeast NADPH-P450 reductase as above so that the concentration of the compound was 0.5 mM. After incubation at 30 °C for 15 hours, the supernatant was obtained by centrifugation (5000 x g, 10 min). Then 2 ml of dichloromethane was added to the supernatant and vigorously stirred and centrifuged (5000 x g, 10 min). The dichloromethane layer was recovered from the separated layer, then evaporated under reduced pressure. The obtained residue was dissolved in 100 µl of acetonitrile, and the solution was analyzed by HPLC under the following conditions.

NADPH and chlorzoxazone were added to a microsomal fraction of yeasts expressing (i) human cytochrome P450 molecular species and yeast NADPH-P450 reductase; or (ii) an artificial fused enzyme comprising human cytochrome P450 molecular species and yeast NADPH-P450 reductase prepared as above so that the concentrations of NADPDH and chlorzoxazone were 0.5 mM and 250  $\mu$ M. Then the

solutions were incubated at 37 °C for 10 min. After that, trichloroacetic acid was added to the solutions so that the concentration of the trichloroacetic acid was about 10% (v/v). Then 2 ml of dichloromethane was added to the solution, and the solution was stirred vigorously and centrifuged (15,000 x g, 5 min). The dichloromethane layer was recovered, and the solvent was removed under reduced pressure. The obtained residue was dissolved in 100  $\mu$ l of acetonitrile and the solution was subjected to analysis by HPLC under the same conditions as above.

All of the yeast cells expressing eleven human P450 molecular species gave hydroxylated chlorzox-azone. P450 2E1 showed high activity, and P450 1A1, P450 1A2, P450 2A6, P450 2D6 showed good activity, while P450 2C8, 2C9, 2B6, 2C18, 2C19 and 3A4 showed moderate activity.

### Ames test using yeast S-9 Mix fraction and microsomal fraction

The Ames test method was in accordance with the customary method described, for example, in Mutat. Res., (1975) 31, 347. 2-Aminoanthrathene which is an arylamine type compound was used as a specimen compound. (1) Rat S-9 Mix supernatant fraction (obtained by homogenizing liver and then subjected to centrifugation (9000 x g, 10 min), manufactured by Kikkoman) containing each kind of rat P450 molecular species at the concentration of 1200 pmol per 1 sample and (2) Yeast S-9 Mix fraction obtained from each kind of yeast cells expressing human P450 or a microsomal fraction prepared from the yeast S-9 Mix fraction were used as a metabolic activation source in the Ames test. As a result, more than 1000 revertant colonies were detected for the compound at 1 µg/plate (90 mm dia.) only in the case of using the yeast S-9 Mix fraction obtained from the yeast cells expressing human P450 1A2 (Saccharomyces cerevisiae AH22/p1A2R) and yeast cells expressing human P450 2E1 (Saccharomyces cerevisiae AH22/p2E1R) and a microsomal fraction prepared from the yeast S-9 Mix fraction, while the amounts of the human P450 molecules present in the Rat S-9 mixture.

The human cytochrome P450 1A2 showed high activity, and human P450 2E1 showed only moderate activity. But the revertant colonies were not found for the human cytochrome P450 3A4, 2C8 and 2A6.

### Metabolism of acetanilide using transformed yeast cells

Acetanilide was added to each of the culture solutions of the transformed yeast cells expressing (i) human cytochrome P450 molecular species and yeast NADPH-P450 reductase; or (ii) an artificial fused enzyme comprising human cytochrome P450 molecular species and yeast NADPH-P450 reductase, so that the concentration of the compound was 5 mM, and the solution was incubated at 30 °C for 15 hours. Then the solution was centrifuged (5000 x g, 10 min) to give a supernatant. The obtained supernatant solution was subjected to the HPLC analysis under the following conditions. The hydroxylated acetanilide was found for all of the tested eleven human P450 molecular species.

Among them, P450 1A2 and 2D6 showed high activity and P450 1A1, 2A6, 2B6, 2C8, 2C9, 2C18, 2C19 and 2E1 showed good activity, while 3A4 showed moderate activity.

#### Conditions for HPLC

Column:

40

45

10

μBondapak C18 (manufactured by Waters Co.)

Carrier:

Methanol:water:acetic acid = 15:84:1

Temperature:

30 · C

Detection:

UV 254 nm

Injection amount: 50 µI

## Metabolism of coumarin using transformed yeast cells

Coumarin was added to 6 ml of each of the culture solutions (SDS synthetic culture medium, cell concentration of about 2.0 x 10<sup>7</sup> cells/ml) of the transformed yeast cells expressing (i) human cytochrome P450 molecular species and yeast NADPH-P450 reductase; or (ii) an artificial fused enzyme comprising human cytochrome P450 molecular species and yeast NADPH-P450 reductase prepared as above, so that the concentration of the compound was 5 mM, and the solution was incubated at 30 °C for 2 or 5 hours. Then the solution was centrifuged (5000 x g, 10 min) to give a supernatant. 62.5 µl of 15% trichloroacetic acid and 2 ml of chloroform were added to the obtained supernatant solution, and the resultant solution was stirred well. The chloroform layer was recovered from the separated layer. Then 4 ml of sodium hydroxide solution containing 0.1 M NaCl was added to the solution and centrifuged again (5000 x g, 10 min). The supernatant fraction was recovered and subjected to fluorescence analysis (ex. 366 nm, em. 452 nm) to

measure the 7-hydroxycoumarin formed. The hydroxylation activity was specifically found only for the yeast cells expressing the human P450 2A6, while other yeast cells showed no activity.

Metabolism of debrisoquine using the microsomal fraction of transformed yeast whole cells

NADPDH and [¹⁴C]debrisoquine were added to each microsomal fraction solution of (i) human cytochrome P450 molecular species and yeast NADPH-P450 reductase; or (ii) an artificial fused enzyme comprising human cytochrome P450 molecular species and yeast NADPH-P450 reductase prepared as above, so that the concentration of the compound was 100 µM and that of NADPH is 6 mM, and the solution was incubated at 30 °C for 30 minutes. Then perchlorate was added to the solution, so that the final concentration of the perchlorate was 10% (v/v). The solution was sufficiently stirred and centrifuged (15,000 x g, 15 min) to give the supernatant. The obtained supernatant was subjected to HPLC analysis according to the following conditions.

Microsomal fractions of yeasts expressing P450 1A1 and 2D6 showed good activity for the hydroxylation of the debrisoquine, while those of yeast cells expressing other human P450 molecular species showed no activity.

Conditions for HPLC

Column:

COSMOSIL 5C18 (manufactured by Nakarai Tesq Co.)

Carrier:

A(acetonitrile)/B(20mM Sodium Perchlorate, pH = 2.5)

20

25

30

Time (minute)	A/B
0-15	9/91
15-30	9/91-25/75 (linear gradient)
30-32	100/0
32-42	9/91

Temperature:

room temperature

Detector:

RI 14 C

Injection amount:

100 µl

#### Metabolism of S-mephenytoin using the microsomal fraction of transformed yeast cells

NADPH and [¹⁴C]S-mephenytoin were added to each microsomal fraction solution of (i) human cytochrome P450 molecular species and yeast NADPH-P450 reductase; or (ii) an artificial fused enzyme comprising human cytochrome P450 molecular species and yeast NADPH-P450 reductase prepared as above, so that the concentration of the compound was 25  $\mu$ M and that of NADPH was 3 mM, and the solution was incubated at 30 °C for 30 minutes. Then the solution was diluted with equal volume of methanol, sufficiently stirred and centrifuged (15,000 x g, 5 min) to give the supernatant. The obtained supernatant was subjected to HPLC analysis according to the following conditions.

Microsomal fractions of yeasts expressing P450 2C19 showed good activity for the hydroxylation of the S-mephenytoin, while those of yeast cells expressing other human P450 molecular species showed no activity.

Conditions for HPLC

Column:

COSMOSIL 5C18 (manufactured by Nakarai Tesq Co.)

Carrier:

A:(Methanol)/(20 mM Potassium phosphate buffer, pH = 7.0) = 40/60

B:Methanol

50

45

Time (minute)	A/B
0-18	100/0
18-20	0/100
20-35	100/0

55

Temperature:

room temperature

Detector:

RI 14 C

Specimen amount:

100 μ1

Results of the hydroxylation activity using human P450 molecular species Human P450 molecular species				numan 7400 molecular species					
	2E1	3A4	1A1	2A6	286	208	2018	2019	2D6
· I	‡	+	++++	‡	+++++++++++++++++++++++++++++++++++++++	+	+	+	‡
		1	1	ı	1	+	+	<b>+</b>	1
	+ + +	+ + +	+ =	1 4	1 4	+ +	ı 4	1 4	1 +
	= ‡	- 1	- *	- 1	- <b>*</b>	- 1	- *	*	*
	: ‡	+	+++	‡	‡	+	<b>+</b>	‡	+++
			,	† † †	•	,	•	•	•
		1	++	,	t	ı	1	,	+ + +
			•	•	ı	,	1	+ + +	•

Metabolism of chlorzoxazone using a mixture of microsomal fractions of transformed yeast cells

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Microsomal fractions of yeast expressing cytochrome P450 prepared as above were mixed in the following molar ratios, and the hydroxylation activities of the mixed solutions were measured using

chlorzoxazone.

_	_		
_	_		
	_		

10

15

P450	System A	System B
3A4	35%	33%
2C9	25%	5.8%
2C8		5.8%
2C18		5.8%
2C19		5.8%
1A2	23%	19%
2E1	17%	15%
1A1		2.4%
2A6		3.0%
2B6		2.4%
2D6		2.4%

The substrate, [\$^4\$C]chlorzoxazone and NADPH were added to the mixed yeast microsomal fractions, so that the concentrations of the compound and NADPH were 382  $\mu$ M and 3 mM. The solutions were incubated at 37 °C for 30 min, and then 1 ml of dichloromethane was added thereto to stop the reaction. After stirring, dichloromethane layer was recovered by centrifugation (10,000 x g, 5 min). Then the solvent was evaporated by the stream of nitrogen gas. The obtained residue was dissolved in 54  $\mu$ l of acetonitrile and 146  $\mu$ l of water, the solution was subjected to HPLC analysis under the following conditions.

Conditions for HPLC

Column:

COSMOSIL 5C18 (manufactured by Nakarai Tesq Co.)

Carrier:

A(Acetonitrile/Water = 27/73)

B(Acetonitrile)

30

35

45

50

Time (minute)	A/B
0-15	100/0
15-17	0/100
17-25	100/0

Temperature:

room temperature

Detector:

RI 14 C

Injection amount:

100 μΙ

The metabolites of chlorzoxazone observed by each of the mixed systems A and B were similar to those metabolites which Guengerich reported based on their experimental results by using human liver microsomal fractions (Guengerich, F.P., Chem. Toxicil., Vol.3, pp.566-573, 1990).

Furthermore, the metabolic turnover numbers were calculated for the human liver microsomal fraction (by Guengerich) and for the present yeast microsomal fractions.

The turnover numbers were calculated to be 1.8 and 1.6 in the mixed systems A and B, respectively. The turnover V for the human liver microsomal fraction was calculated using  $V_{max}$ ,  $K_m$  and substrate concentration [S] described in the literature according to the following manner. The results are shown in Table 2. The values somewhat varied due to the difference of individuals, the lowest value being 1.0 and the highest value being 5.9. The values of V for the mixed system B and A fell within this range, both of which were the same level. It was confirmed that the four kinds of molecular species in system A can well reproduce the metabolic system in human liver in vitro.

A turnover V for human cytochrome P450 at an optional substrate concentration can be calculated by substituting  $V_{max}$  and  $K_m$  described in the literature and substrate concentration [S] of the present example into the Michaelis-Menten's equation:

$$V = (V_{max} * [S])/(K_m + [S])$$

5

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15

20

25

30

35

40

Table 2

Liver sample	Metabolic turnover V [product nmol/nmo P450/min]
#1001	5.9
KDL 14	2.2
KDL 21	1.7
KDL 23	3.0
KDL 27	5.0
H 10	1.1
H 11	1.0
H 12	4.2
H 13	3.3
H 14	2.1
H 15	4.3
H 16	4.0
H 17	3.6
H 18	3.4

Metabolism of debrisoquine using mixture of microsomal fractions of transformed yeast cells

Microsomal fractions of yeasts expressing human cytochrome P450 were mixed, and the hydroxylation activity of the mixed fraction was measured using debrisoquine. The mixing molar ratio of the human cytochrome P450 molecular species were as follows:

Molar ratio
33%
5.8%
5.8%
5.8%
5.8%
19%
15%
2.4%
2.4%
2.4%

The substrate debrisoquine and NADPH were added to the mixed microsomal fraction solutions, so that the concentrations were 100  $\mu$ M for the NADPH and 6 mM for the compound. After the mixture was incubated at 37 °C for 30 min, 50  $\mu$ l of 60% perchlorate was added to the solution to stop the reaction. The concentration of the perchlorate was finally 12.5% (v/v). After vigorous stirring, the mixture was centrifuged (15,000 x g, 5 min) to recover the supernatant, which was subjected to HPLC analysis under the same conditions used for analyzing the metabolites of debrisoquine.

The metabolites well coincided with the metabolites which Kronbach reported based on the experiments to metabolize the debrisoquine using the human liver microsome (Methods in Enzymology, Vol.206, pp.509-517, 1991).

### Metabolism of S-mephenytoin using mixture of microsomal fractions of transformed yeast cells

Microsomal fractions of yeasts expressing various human cytochrome P450 prepared were mixed, and the hydroxylation activity of the mixed fraction was measured for S-mephenytoin. The mixing ratio of the human cytochrome P450 molecular species was the same as that of the mixing system B as described above.

The substrate, [14 C]S-mephenytoin and NADPH were added to the mixed microsomal fraction solutions, so that the concentrations were 28  $\mu$ M for the NADPH and 6 mM for the compound. After the mixture was incubated at 37 °C for 30 min, 250  $\mu$ l of methanol was added to the solution to stop the reaction. After vigorous stirring, the mixture was centrifuged (15,000 x g, 5 min) to recover the supernatant, which was subjected to HPLC analysis under the same conditions used for the hydroxylation of S-mephenytoin using microsomal fraction. The metabolites obtained well coincided with the metabolites which Goldstein reported based on the experiments to metabolize the S-mephenytoin using the human liver microsome (Biochemistry, Vol.33, pp.1743-1752, 1994).

10 SEQUENCE LISTING (1) GENERAL INFORMATION: (i) APPLICANT: 15 (A) NAME: Sumitomo Chemical Company, Limited (B) STREET: 5-33, Kitahama 4-chome, Chuo-ku, (C) CITY: Osaka-shi, Osaka-fu (E) COUNTRY: Japan (F) POSTAL CODE (ZIP): none 20 (ii) TITLE OF INVENTION: METHOD FOR SAFETY EVALUATION OF CHEMICAL COMPOUND USING RECOMBINANT YEAST EXPRESSING HUMAN CYTOCHROME P450 (iii) NUMBER OF SEQUENCES: 61 25 (iv) COMPUTER READABLE FORM: (A) MEDIUM TYPE: Floppy disk (B) COMPUTER: IBM PC compatible (C) OPERATING SYSTEM: PC-DOS/MS-DOS (D) SOFTWARE: PatentIn Release #1.0, Version #1.25 (EPO) 30 (vi) PRIOR APPLICATION DATA: (A) APPLICATION NUMBER: JP 201120/1993 (B) FILING DATE: 20-JUL-1993 (vi) PRIOR APPLICATION DATA: (A) APPLICATION NUMBER: JP 180246/1993 35 (B) FILING DATE: 21-JUL-1993 (vi) PRIOR APPLICATION DATA: (A) APPLICATION NUMBER: JP 208279/1993 (B) FILING DATE: 30-JUL-1993 40 (2) INFORMATION FOR SEQ ID NO: 1: (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 1551 base pairs (B) TYPE: nucleic acid 45 (C) STRANDEDNESS: double (D) TOPOLOGY: linear (ix) FEATURE: (A) NAME/KEY: CDS 50 (B) LOCATION: 1..1548 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 1: 48 ATG GCA TTG TCC CAG TCT GTT CCC TTC TCG GCC ACA GAG CTC CTC CTG 55 Met Ala Leu Ser Gln Ser Val Pro Phe Ser Ala Thr Glu Leu Leu

1

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	GCC Ala	TCT Ser	GCC Ala	ATC Ile 20	TTC Phe	TGC Cys	CTG Leu	GTA Val	TTC Phe 25	TGG Trp	GTG Val	CTC Leu	AAG Lys	GGT Gly 30	TTG Leu	AGG Arg	96
5	CCT Pro	CGG Arg	GTC Val 35	CCC Pro	AAA Lys	GGC Gly	CTG Leu	AAA Lys 40	AGT Ser	CCA Pro	CCA Pro	GAG Glu	CCA Pro 45	TGG Trp	GGC Gly	TGG Trp	144
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	CTG Leu 65	TCA Ser	AGG Arg	ATG Met	AGC Ser	CAG Gln 70	CGC Arg	TAC Tyr	GGG Gly	GAC Asp	GTC Val 75	CTG Leu	CAG Gln	ATC Ile	CGC Arg	ATT Ile 80	240
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20	GCC Ala	CTG Leu	GTG Val	CGG Arg 100	CAG Gln	GGC Gly	GAC Asp	GAT Asp	TTC Phe 105	AAG Lys	GGC Gly	CGG Arg	CCT Pro	GAC Asp 110	CTC Leu	TAC Tyr	336
	ACC Thr	TCC Ser	ACC Thr 115	CTC Leu	ATC Ile	ACT Thr	GAT Asp	GGC Gly 120	CAG Gln	AGC Ser	TTG Leu	ACC Thr	TTC Phe 125	AGC Ser	ACA Thr	GAC Asp	384
25	TCT Ser	GGA Gly 130	CCG Pro	GTG Val	TGG Trp	GCT Ala	GCC Ala 135	CGC Arg	CGG <b>Ar</b> g	CGC <b>Arg</b>	CTG Leu	GCC Ala 140	CAG Gln	AAT Asn	GCC Ala	CTC Leu	432
30	AAC Asn 145	ACC Thr	TTC Phe	TCC Ser	ATC Ile	GCC Ala 150	TCT Ser	GAC Asp	CCA Pro	GCT Ala	TCC Ser 155	TCA Ser	TCC Ser	TCC Ser	TGC Cys	TAC Tyr 160	480
	CTG Leu	GAG Glu	GAG Glu	CAT His	GTG Val 165	AGC Ser	AAG Lys	GAG Glu	GCT Ala	AAG Lys 170	GCC Ala	CTG Leu	ATC Ile	AGC Ser	AGG Arg 175	TTG Leu	528
35	CAG Gln	G <b>A</b> G Glu	CTG Leu	ATG Met 180	GCA Ala	GGG Gly	CCT Pro	GGG Gly	CAC His 185	TTC Phe	GAC Asp	CCT Pro	TAC Tyr	AAT Asn 190	CAG Gln	GTG Val	576
40	GTG Val	GTG Val	TCA Ser 195	GTG Val	GCC Ala	AAC Asn	GTC Val	ATT Ile 200	GGT Gly	GCC Ala	ATG Met	TGC Cys	TTC Phe 205	GGA Gly	CAG Gln	CAC His	624
	Phe	Pro	Glu	Ser	Ser	Asp	Glu	Met	Leu	Ser	Leu	GTG Val 220	Lys	AAC Asn	ACT Thr	CAT His	672
<b>4</b> 5	GAG Glu 225	TTC Phe	GTG Val	GAG Glu	ACT Thr	GCC Ala 230	TCC Ser	TCC Ser	GGG Gly	AAC Asn	CCC Pro 235	CTG Leu	GAC Asp	TTC Phe	TTC Phe	CCC Pro 240	720
	ATC Ile	CTT Leu	CGC <b>A</b> rg	TAC Tyr	CTG Leu 245	CCT Pro	AAC Asn	CCT Pro	GCC Ala	CTG Leu 250	CAG Gln	AGG Arg	TTC Phe	AAG Lys	GCC Ala 255	TTC Phe	768

	AAC Asn	CAG Gln	AGG Arg	TTC Phe 260	CTG Leu	TGG Trp	TTC Phe	CTG Leu	CAG Gln 265	AAA Lys	ACA Thr	GTC Val	CAG Gln	GAG Glu 270	CAC His	TAT Tyr	81	.6
5	CAG Gln	GAC Asp	TTT Phe 275	GAC Asp	AAG Lys	AAC Asn	AGT Ser	GTC Val 280	CGG Arg	GAC Asp	ATC Ile	ACG Thr	GGT Gly 285	GCC Ala	CTG Leu	TTC Phe	86	<b>4</b>
10		CAC His 290															91	.2
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	AGG Arg	GAG Glu	CGG Arg 355	CGG Arg	CCC Pro	CGG Arg	CTC Leu	TCT Ser 360	GAC Asp	AGA Arg	CCC Pro	CAG Gln	CTG Leu 365	CCC Pro	TAC Tyr	TTG Leu	110	4
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30	ACC Thr 385	ATC Ile	CCC Pro	CAC His	AGC Ser	ACA Thr 390	ACA Thr	AGG Arg	GAC Asp	ACA Thr	ACG Thr 395	CTG Leu	AAT Asn	GGC Gly	TTC Phe	TAC Tyr 400	120	0 (
30	ATC Ile	CCC Pro	AAG Lys	AAA Lys	TGC Cys 405	TGT Cys	GTC Val	TTC Phe	GTA Val	AAC Asn 410	CAG Gln	TGG Trp	CAG Gln	GTC Val	AAC Asn 415	CAT His	124	. 8
35	GAC Asp	CCA Pro	GAG Glu	CTG Leu 420	TGG Trp	GAG Glu	GAC Asp	CCC Pro	TCT Ser 425	GAG Glu	TTC Phe	CGG <b>A</b> rg	CCT Pro	GAG Glu 430	CGG <b>A</b> rg	TTC Phe	129	16
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<b>4</b> 5	AAG Lys 465	TGG Trp	GAG Glu	ATC Ile	TTC Phe	CTC Leu 470	TTC Phe	CTG Leu	GCC Ala	ATC Ile	CTG Leu 475	CTA Leu	CAG Gln	CAA Gln	CTG Leu	GAG Glu 480	144	10

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5	CTG Leu	ACC Thr	ATG Met	AAG Lys 500	CAC His	GCC Ala	CGC Arg	TGT Cys	GAA Glu 505	CAT His	GTC Val	CAG Gln	GCG Ala	CGG Arg 510	CTG Leu	CGC Arg	1536
10			ATC Ile 515		TGA												1551
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15		•	( <i>1</i>	A) LE 3) TY	ENGTI (PE :	I: 51 amir	RACTE 16 and 10 ac 1ine	mino cid									
		•					prot										
20	Mak						(PTIC						Glu	Leu	Leu	Leu	
	1				5					10					15		
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5

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	Glu 225	Phe	Val	Glu	Thr	Ala 230	Ser	Ser	Gly	Asn	Pro 235	Leu	Asp	Phe	Phe	Pro 240
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	Pro	Glu	Ile	Gln 340	Arg	Lys	Ile	Gln	Lys 345	Glu	Leu	Asp	Thr	<b>V</b> al 350	Ile	Gly
30	Arg	Glu	Arg 355	Arg	Pro	Arg	Leu	Ser 360	Asp	Arg	Pro	Gln	Leu 365	Pro	Tyr	Leu
	Glu	Ala 370	Phe	Ile	Leu	Glu	Thr 375	Phe	Arg	His	Ser	Ser 380	Phe	Leu	Pro	Phe
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5	Phe	Ser	Ile 515	Asn													
	(2)	INF	OR <b>MA</b> '	r'ION	FOR	SEQ	ID I	<b>N</b> O:	3:								
10		(i)	() () ()	A) L1 B) T C) S	ENGTI YPE : I'R <b>AN</b> I	H: 14 nuc DEDNI	CTER 473   leic ESS: line	oase acie doul	pai: d	rs							
15		(ix)	()	ATURI A) NA B) L	AME/		CDS	1470									
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	CCC Pro	ACT Thr	CCT Pro 35	CTC Leu	CCA Pro	GTG Val	ATT Ile	GGA Gly 40	AAT Asn	ATC Ile	CTA Leu	CAG Gln	ATA Ile 45	GGT Gly	ATT Ile	AAG Lys	144
30									CTC Leu								192
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33	GAA Glu	GCA Ala	GTG Val	AAG Lys	GAA Glu 85	GCC Ala	CTG Leu	ATT Ile	GAT Asp	CTT Leu 90	GGA Gly	GAG Glu	GAG Glu	TTT Phe	TCT Ser 95	GGA Gly	288
40	AGA Arg	GGC Gly	ATT Ile	TTC Phe 100	CCA Pro	CTG Leu	GCT Ala	GAA Glu	AGA Arg 105	GCT Ala	AAC Asn	AGA Arg	GGA Gly	TTT Phe 110	GGA Gly	ATT Ile	336
	GTT Val	TTC Phe	AGC Ser 115	AAT Asn	GGA Gly	AAG Lys	AAA Lys	TGG Trp 120	AAG Lys	G <b>A</b> G Glu	ATC Ile	CGG Arg	CGT Arg 125	TTC Phe	TCC Ser	CTC Leu	384
<b>4</b> 5	ATG Met	ACG Thr 130	CTG Leu	CGG Arg	AAT Asn	TTT Phe	GGG Gly 135	ATG Met	GGG Gly	AAG Lys	AGG Arg	AGC Ser 140	ATT Ile	GAG Glu	GAC Asp	CGT Arg	432
50																	

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5	GCC Ala	TCA Ser	CCC Pro	TGT Cys	GAT Asp 165	CCC Pro	ACT Thr	TTC Phe	ATC Ile	CTG Leu 170	GGC Gly	TGT Cys	GCT Ala	CCC Pro	TGC Cys 175	AAT Asn	528
10	GTG Val	ATC Ile	TGC Cys	TCC Ser 180	ATT Ile	ATT Ile	TTC Phe	CAT His	AAA Lys 185	CGT Arg	TTT Phe	GAT Asp	TAT Tyr	AAA Lys 190	GAT Asp	CAG Gln	576
	CAA Gln	TTT Phe	CTT Leu 195	AAC Asn	TTA Leu	ATG Met	GAA Glu	AAG Lys 200	TTG Leu	AAT Asn	GAA Glu	AAC Asn	ATC Ile 205	AAG Lys	ATT Ile	TTG Leu	624
15	AGC Ser	AGC Ser 210	CCC Pro	TGG Trp	ATC Ile	CAG Gln	ATC Ile 215	TGC Cys	AAT Asn	AAT Asn	TTT Phe	TCT Ser 220	CCT Pro	ATC Ile	ATT Ile	GAT Asp	672
20	TAC Tyr 225	TTC Phe	CCG Pro	GGA Gly	ACT Thr	CAC His 230	AAC Asn	AAA Lys	TTA Leu	CTT Leu	AAA Lys 235	AAC Asn	GTT Val	GCT Ala	TTT Phe	ATG Met 240	720
	AAA <b>Ly</b> s	AGT Ser	TAT Tyr	ATT Ile	TTG Leu 245	G <b>AA</b> Glu	AAA Lys	GTA Val	Lys Lys	GAA Glu 250	CAC His	CAA Gln	GAA Glu	TCA Ser	ATG Met 255	GAC Asp	768
25	ATG Met	AAC Asn	AAC Asn	CCT Pro 260	CAG Gln	GAC Asp	TTT Phe	ATT Ile	GAT Asp 265	TGC Cys	TTC Phe	CTG Leu	ATG Met	AAA Lys 270	ATG Met	GAG Glu	816
30	AAG Lys	GAA Glu	AAG Lys 275	CAC His	AAC Asn	CAA Gln	CCA Pro	TCT Ser 280	GAA Glu	TTT Phe	ACT Thr	ATT Ile	GAA Glu 285	AGC Ser	TTG Leu	GAA Glu	864
	AAC Asn	ACT Thr 290	GCA Ala	GTT Val	GAC Asp	TTG Leu	TTT Phe 295	GGA Gly	GCT Ala	GGG Gly	ACA Thr	GAG Glu 300	ACG Thr	ACA Thr	AGC Ser	ACA Thr	912
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40	GCT Ala	AAA Lys	GTC Val	CAG Gln	GAA Glu 325	G <b>A</b> G Glu	ATT Ile	GAA Glu	CGT Arg	GTG Val 330	ATT Ile	GGC Gly	AGA Arg	AAC Asn	CGG Arg 335	AGC Ser	1008
	CCC Pro	TGC Cys	ATG Met	CAA Gln 340	GAC Asp	AGG Arg	AGC Ser	CAC His	ATG Met 345	Pro	TAC Tyr	Thr	GAT Asp	Ala	Val	GTG Val	1056
<b>4</b> 5	CAC His	GAG Glu	GTC Val 355	CAG Gln	AGA Arg	TAC Tyr	ATT Ile	GAC Asp 360	Leu	CTC Leu	CCC Pro	ACC Thr	AGC Ser 365	CTG Leu	CCC Pro	CAT His	1104
	GCA Ala	GTG Val 370	Thr	TGT Cys	GAC Asp	ATT Ile	AAA Lys 375	Phe	AGA Arg	AAC Asn	TAT Tyr	CTC Leu 380	Ile	CCC Pro	AAG Lys	GGC Gly	1152

	ACA Thr 385	ACC Thr	ATA Ile	TTA Leu	ATT Ile	TCC Ser 390	CTG Leu	ACT Thr	TCT Ser	GTG Val	CTA Leu 395	CAT His	GAC Asp	AAC Asn	AAA Lys	GAA Glu 400	1200
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	CGG Arg	ATT Ile	TGT Cys 435	GTG Val	GGA Gly	GAA Glu	GCC Ala	CTG Leu 440	GCC Ala	GGC Gly	ATG Met	GAG Glu	CTG Leu 445	TTT Phe	TTA Leu	TTC Phe	1344
15	CTG Leu	ACC Thr 450	TCC Ser	ATT Ile	TTA Leu	CAG Gln	AAC Asn 455	TTT Phe	AAC Asn	CTG Leu	rys Y <b>yy</b>	TCT Ser 460	CTG Leu	GTT Val	GAC Asp	CCA Pro	1392
20	AAG Lys 465	AAC Asn	CTT Leu	GAC Asp	ACC Thr	ACT Thr 470	CCA Pro	GTT Val	GTC Val	AAT Asn	GGA Gly 475	TTT Phe	GCC <b>A</b> la	TCT Ser	GTG Val	CCG Pro 480	1440
	CCC Pro	TTC Phe	TAC Tyr	CAG Gln	CTG Leu 485	TGC Cys	TTC Phe	ATT Ile	CCT Pro	GTC Val 490	TGA						1473
25	(2)			CION SEQUE													
30		,	( <i>I</i>	A) LE B) TY D) TO	ENGTI (PE :	I: 49 amir	90 an	nino cid	acio	ls							
				ECUI													
		(xi)	SEC	QUENC	CE DI	ESCR	PTIC	ON: S	SEQ I	D NO	): <b>4</b> :	:				•	
35	Met 1	Asp	Ser	Ile	Val 5	Ser	Leu	Val	Leu	Cys 10	Leu	Ser	Суз	Leu	Leu 15	Leu	
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	Asp	Ile 50	Ser	Lys	Ser	Leu	Thr 55	Asn	Leu	Ser	Lys	Val 60	Tyr	Gly	Pro	Val	
<b>4</b> 5	Phe 65	Thr	Leu	Tyr	Phe	Gly 70	Leu	Lys	Pro	Ile	Val 75	Val	Leu	His	Gly	Tyr 80	
	Glu	Ala	Val	Lys	Glu 85	Ala	Leu	Ile	Asp	Leu 90	Gly	Glu	Glu	Phe	Ser 95	Gly	

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	Ala	Ser	Pro	Cys	Asp 165	Pro	Thr	Phe	Ile	Leu 170	Gly	Суѕ	Ala	Pro	Cys 175	Asn
15	Val	Ile	Суѕ	Ser 180	Ile	Ile	Phe	His	Lys 185	Arg	Phe	Asp	Tyr	Lys 190	Asp	Gln
	Gln	Phe	Leu 195	Asn	Leu	Met	Glu	L <b>y</b> s 200	Leu	Asn	Glu	Asn	Ile 205	Lys	Ile	Leu
20	Ser	Ser 210	Pro	Trp	Ile	Gln	Ile 215	Суѕ	Asn	Asn	Phe	Ser 220	Pro	Ile	Ile	Asp
	Tyr 225	Phe	Pro	Gly	Thr	His 230	Asn	Lys	Leu	Leu	Lys 235	Asn	Val	Ala	Phe	Met 240
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	Met	Asn	Asn	Pro 260	Gln	Asp	Phe	Ile	Asp 265	Cys	Phe	Leu	Met	Lys 270	Met	Glu
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40	Pro	Cys	Met	Gln 340	Ąsp	Arg	Ser	His	Met 345	Pro	Tyr	Thr	Asp	<b>Ala</b> 350	Val	Val
	His	Glu	Val 355	Gln	Arg	Tyr	Ile	Asp 360	Leu	Leu	Pro	Thr	Ser 365	Leu	Pro	His
<b>4</b> 5	Ala	Val 370	Thr	Cys	Asp	Ile	<b>Lys</b> 375	Phe	Arg	Asn	Tyr	Leu 380	Ile	Pro	Lys	Gly
	Thr 385	Thr	Ile	Leu	Ile	Ser 390	Leu	Thr	Ser	Val	Leu 395	His	Asp	Asn	Lys	Glu <b>4</b> 00
50	Phe	Pro	Asn	Pro	Glu 405	Met	Phe	Asp	Pro	His 410	His	Phe	Leu	Asp	Glu 415	Gly

	Gly	Asn	Phe	Lys 420	Lys	Ser	Lys	Tyr	Phe 425	Met	Pro	Phe	Ser	Ala 430	Gly	Lys		
5	Arg	Ile	Cys 435	Val	Gly	Glu	Ala	Leu 440	Ala	Gly	Met	Glu	Leu 445	Phe	Leu	Phe		
	Leu	Thr 450	Ser	Ile	Leu	Gln	Asn 455	Phe	Asn	Leu	Lys	Ser 460	Leu	Val	Asp	Pro		
10	Lys 465	Asn	Leu	Asp	Thr	Thr 470	Pro	Val	Val	Asn	Gly 475	Phe	Ala	Ser	Val	Pro 480		
	Pro	Phe	Tyr	Gln	Leu 485	Cys	Phe	Ile	Pro	Val 490								
15	(2)	(i)	SEQ ( <i>F</i> (E	OUENC A) LE B) TY C) ST	CE CH ENGTH (PE: TRANI	HARAC H: 14 nucl	ID N CTERI 182 k leic ESS: line	STIC ase acid doub	CS: pain	rs								
20			( <i>P</i>	3) LC	ME/I		11											
		(xi)	SEÇ	QUENC	CE DE	ESCRI	PTIC	ON: 5	SEQ 1	ID NO	D: 5:	:						
	ATG	TCT Ser	GCC Ala	CTC Leu	GGA Gly 5	GTC Val	ACC Thr	GTG Val	GCC Ala	CTG Leu 10	CTG Leu	GTG Val	TGG Trp	GCG <b>A</b> la	GCC Ala 15	TTC Phe		48
30		CTG Leu																96
35		CCA Pro														TTG Leu	:	144
	GAA Glu	TTG Leu 50	AAG Lys	TAA Asn	ATT Ile	CCC Pro	AAG Lys 55	TCC Ser	TTC Phe	ACC Thr	CGG Arg	TTG Leu 60	GCC Ala	CAG Gln	CGC Arg	TTC Phe	:	192
<b>4</b> 0	GGG Gly 65	CCG Pro	GTG Val	TTC Phe	ACG Thr	CTG Leu 70	TAC Tyr	GTG Val	GGC Gly	TCG Ser	CAG Gln 75	CGC Arg	ATG Met	GTG Val	GTG Val	ATG Met 80	:	240
<b>4</b> 5	CAC His	GGC Gly	TAC Tyr	<b>AA</b> G Lys	GCG Ala 85	GTG Val	<b>AA</b> G Lys	G <b>AA</b> Glu	GCG Ala	CTG Leu 90	CTG Leu	GAC Asp	TAC 'Fyr	AAG Lys	GAC Asp 95	GAG Glu	:	288
		TCG Ser															:	336

	GGA Gly	ATC Ile	ATT Ile 115	TTT Phe	AAT Asn	AAT Asn	GGA Gly	CCT Pro 120	ACC Thr	TGG Trp	<b>AA</b> G Lys	GAC Asp	ATC Ile 125	CGG Arg	CGG Arg	TTT Phe	384
5	TCC Ser	CTG Leu 130	ACC Thr	ACC Thr	CTC Leu	CGG Arg	AAC Asn 135	TAT Tyr	GGG Gly	ATG Met	GGG Gly	AAA Lys 140	CAG Gln	GGC Gly	AAT Asn	GAG Glu	432
10	AGC Ser 145	CGG Arg	ATC Ile	CAG Gln	AGG Arg	GAG Glu 150	GCC Ala	CAC His	T <b>T</b> C Phe	CTG Leu	CTG Leu 155	GAA Glu	GCA Ala	CTC Leu	AGG Arg	AAG Lys 160	480
	ACC Thr	CAA Gln	GGC Gly	CAG Gln	CCT Pro 165	TTC Phe	GAC Asp	CCC Pro	ACC Thr	TTC Phe 170	CTC Leu	ATC Ile	GGG Gly	TGC Cys	GCG Ala 175	CCC Pro	5 <b>28</b>
15	TGC Cys	AAC Asn	GTC Val	ATA Ile 180	GCC Ala	GAC Asp	ATC Ile	CTC Leu	TTC Phe 185	CGC Arg	AAG Lys	CAT His	TTT Phe	GAC Asp 190	TAC Tyr	AAT Asn	5 <b>7</b> 6
20	GAT Asp	GAG Glu	AAG Lys 195	TTT Phe	CTA Leu	AGG Arg	CTG Leu	ATG Met 200	TAT Tyr	TTG Leu	TTT Phe	AAT Asn	GAG Glu 205	AAC Asn	TTC Phe	CAC His	624
	CTA Leu	CTC Leu 210	AGC Ser	ACT Thr	CCC Pro	TGG Trp	CTC Leu 215	CAG Gln	CTT Leu	TAC Tyr	AAT Asn	AAT Asn 220	TTT Phe	CCC Pro	AGC Ser	TTT Phe	672
25	CTA Leu 225	CAC His	TAC Tyr	TTG Leu	CCT Pro	GGA Gly 230	AGC Ser	CAC His	AGA Arg	AAA Lys	GTC Val 235	ATA Ile	AAA Lys	AAT Asn	GTG Val	GCT Ala 240	720
30	GAA Glu	GTA Val	AAA Lys	G <b>A</b> G Glu	TAT Tyr 245	GTG Val	TCT Ser	GAA Glu	AGG Arg	GTG Val 250	AAG Lys	GAG Glu	CAC	CAT His	CAA Gln 255	TCT Ser	768
	CTG Leu	GAC Asp	CCC Pro	AAC Asn 260	TGT Cys	CCC Pro	CGG Arg	GAC Asp	CTC Leu 265	ACC Thr	GAC Asp	TGC Cys	CTG Leu	CTC Leu 270	GTG Val	GAA Glu	816
<i>3</i> 5	ATG Met	GAG Glu	AAG Lys 275	G <b>AA</b> Glu	AAG Lys	CAC His	AGT Ser	GCA Ala 280	GAG Glu	CGC Arg	TTG Leu	TAC Tyr	ACA Thr 285	ATG Met	GAC Asp	GGT Gly	864
40	ATC Ile	ACC Thr 290	GTG Val	ACT Thr	GTG Val	GCC Ala	GAC Asp 295	CTG Leu	TTC Phe	TTT Phe	GCG Ala	GGG Gly 300	ACA Thr	GAG Glu	ACC Thr	ACC Thr	912
	AGC Ser 305	Thr	ACT Thr	CTG Leu	AGA Arg	TAT Tyr 310	Gly	CTC Leu	CTG Leu	ATT Ile	CTC Leu 315	ATG Met	AAA Lys	TAC Tyr	CCT Pro	GAG Glu 320	960
<b>4</b> 5	ATC Ile	GAA Glu	GAG Glu	AAG Lys	CTC Leu 325	His	GAA Glu	G <b>A</b> A Glu	ATT	GAC Asp 330	Arg	GTG Val	ATT Ile	GGG Gly	CCA Pro 335	AGC Ser	1008

	CGA Arg	ATC Ile	CCT Pro	GCC Ala 340	ATC Ile	AAG Lys	GAT Asp	AGG Arg	CAA Gln 345	GAG Glu	ATG Met	CCC Pro	TAC Tyr	ATG Met 350	GAT Asp	GCT Ala	1056
5	GTG Val	GTG Val	CAT His 355	GAG Glu	ATT Ile	CAG Gln	CGG Arg	TTC Phe 360	ATC Ile	ACC Thr	CTC Leu	GTG Val	CCC Pro 365	TCC Ser	AAC Asn	CTG Leu	1104
10	CCC Pro	CAT His 370	GAA Glu	GCA Ala	ACC Thr	CGA Arg	GAC Asp 375	ACC Thr	ATT Ile	TTC Phe	AGA Arg	GGA Gly 380	TAC Tyr	CTC Leu	ATC Ile	CCC Pro	1152
	AAG Lys 385	GGC Gly	ACA Thr	GTC Val	GTA Val	GTG Val 390	CCA Pro	ACT Thr	CTG Leu	GAC Asp	TCT Ser 395	GTT Val	TTG Leu	TAT Tyr	GAC Asp	AAC Asn 400	1200
15	CAA Gln	GAA Glu	TTT Phe	CCT Pro	GAT Asp 405	CCA Pro	GAA Glu	AAG Lys	TTT Phe	AAG Lys 410	CCA Pro	GAA Glu	CAC His	TTC Phe	CTG Leu 415	AAT Asn	1248
20	G <b>AA</b> Glu	AAT Asn	GGA Gly	AAG Lys 420	TTC Phe	AAG Lys	TAC Tyr	AGT Ser	GAC Asp 425	TAT Tyr	TTC Phe	AAG Lys	CCA Pro	TTT Phe 430	TCC Ser	ACA Thr	1296
	GGA Gly	AAA Lys	CGA Arg 435	GTG Val	TGT Cys	GCT Ala	GGA Gly	GAA Glu 440	GGC Gly	CTG Leu	GCT Ala	CGC Arg	ATG Met 445	GAG Glu	TTG Leu	TTT Phe	1344
25	CTT Leu	TTG Leu 450	TTG Leu	TGT Cys	GCC Ala	ATT Ile	TTG Leu 455	CAG Gln	CAT His	TTT Phe	AAT Asn	TTG Leu 460	AAG Lys	CCT Pro	CTC Leu	GTT Val	1392
30	GAC <b>As</b> p 465	CCA Pro	AAG Lys	GAT Asp	ATC Ile	GAC Asp 470	CTC Leu	AGC Ser	CCT Pro	ATA Ile	CAT His 475	ATT Ile	GGG Gly	TTT Phe	GGC Gly	TGT Cys 480	1440
	ATC Ile	CCA Pro	CCA Pro	CGT Arg	TAC Tyr 485	AAA Lys	CTC Leu	TGT Cys	GTC Val	ATT Ile 490	CCC Pro	CGC Arg	TCA Ser	TGA			1482
35	(2)	INF	ORMA'	TION	FOR	SEQ	ID I	<b>1</b> 0: (	<b>5</b> :								
40			() ()	SEQUI A) LI B) T D) T	ENGTI YPE :	H: 49 amir	93 ar	mino cid									
				LECU			_										
				QUEN												ph	
<b>4</b> 5	Met 1	Ser	Ala	Leu	Gly 5	Val	Thr	Val	Ala	Leu 10	Leu	Val	Trp	Ala	Ala 15	Phe	

Leu Leu Leu Val Ser Met Trp Arg Gln Val His Ser Ser Trp Asn Leu 20 25 30

	Pro	Pro	Gly 35	Pro	Phe	Pro	Leu	Pro 40	Ile	Ile	Gly	Asn	Leu 45	Phe	Gln	Leu
5	Glu	Leu 50	Lys	Asn	Ile	Pro	Lys 55	Ser	Phe	Thr	Arg	Leu 60	Ala	Gln	Arg	Phe
	Gly 65	Pro	Val	Phe	Thr	Leu 70	Tyr	Val	Gly	Ser	Gln 75	Arg	Met	Val	Val	Met 80
10	His	Gly	Tyr	Lys	Ala 85	Val	Lys	Glu	Ala	Leu 90	Leu	Asp	Tyr	Lys	<b>As</b> p 95	Glu
	Phe	Ser	Gly	Arg 100	Gly	Asp	Leu	Pro	<b>A</b> la 105	Phe	His	Ala	His	Arg 110	Asp	Arg
15	Gly	Ile	Ile 115	Phe	Asn	naA	Gly	Pro 120	Thr	Trp	Lys	Asp	Ile 125	Arg	Arg	Phe
	Ser	Leu 130	Thr	Thr	Leu	Arg	Asn 135	Tyr	Gly	Met	Gly	Lys 140	Gln	Gly	Asn	Glu
20	145					150					155			Leu		160
					165					170				Cys	175	
25	-			180					185					Asp 190		
	-		195					200					205	Asn		
30		210					215					220		Pro		
	225		-			230					235			Asn		240
<b>3</b> 5					245					250				His	255	
		_		260					265					Leu 270		
40	Met	Glu	<b>Lys</b> 275	Glu	Lys	His	Ser	Ala 280	Glu	Arg	Leu	Tyr	Thr 285	Met	Asp	Gly
		290					295					300		Glu		
<b>4</b> 5	305					310					315			Tyr		320
	Ile	Glu	Glu	Lys	Leu 325	His	Glu	Glu	Ile	Asp 330	Arg	Val	Ile	Gly	Pro 335	Ser
50	Arg	Ile	Pro	Ala 340	Ile	Lys	Asp	Arg	Gln 345		Met	Pro	Tyr	Met 350	Asp	Ala

	Val	Val	His 355	Glu	Ile	Gln	Arg	Phe 360	Ile	Thr	Leu	Val	Pro 365	Ser	Asn	Leu	
5	Pro	His 370	Glu	Ala	Thr	Arg	Asp 375	Thr	Ile	Phe	Arg	Gly 380	Tyr	Leu	Ile	Pro	
	Lys 385	Gly	Thr	Val	Val	Val 390	Pro	Thr	Leu	Asp	Ser 395	Val	Leu	Tyr	Asp	Asn 400	
10	Gln	Glu	Phe	Pro	Asp 405	Pro	Glu	Lys	Phe	Lys 410	Pro	Glu	His	Phe	Leu <b>4</b> 15	Asn	
	Glu	Asn	Gly	Lys 420	Phe	Lys	Tyr	Ser	Asp 425	Tyr	Phe	Lys	Pro	Phe 430	Ser	Thr	
15	Gly	Lys	<b>A</b> rg <b>4</b> 35	Val	Cys	Ala	Gly	Glu <b>44</b> 0	Gly	Leu	Ala	Arg	Met 445	Glu	Leu	Phe	
	Leu	Leu 450	Leu	Cys	Ala	Ile	Leu 455	Gln	His	Phe	Asn	Leu 460	Lys	Pro	Leu	Val	
20	Asp 465	Pro	Lys	Asp	Ile	Asp 470	Leu	Ser	Pro	Ile	His 475	Ile	Gly	Phe	Gly	Cys 480	
	Ile	Pro	Pro	Arg	Tyr 485	Lys	Leu	Суѕ	Val	Ile 490	Pro	Arg	Ser				
25	(2)		SEQ ( <i>P</i> (E	OUENC (A) LE (B) TY (C) ST	CE CH ENGTH PE: TRANI	SEQ HARACH: 15 nucl DEDNE OGY:	CTERI 512 k .eic ESS:	STIC ase acid doub	CS: pair	aî							
30		(ix)	(7		ME/F	KEY:		.509									
35						ESCR											
	ATG Met 1	GCT Ala	CTC Leu	ATC Ile	CCA Pro 5	GAC Asp	TTG Leu	GCC Ala	ATG Met	GAA Glu 10	ACC Thr	TGG Trp	CTT Leu	CTC Leu	CTG Leu 15	GCT Ala	4.8
40	GTC <b>Va</b> l	AGC Ser	CTG Leu	GTG Val 20	CTC Leu	CTC Leu	TAT Tyr	CTA Leu	TAT Tyr 25	GGA Gly	ACC Thr	CAT His	TCA Ser	CAT His 30	GGA Gly	CTT Leu	96
<b>4</b> 5	TTT Phe	AAG Lys	AAG Lys 35	CTT Leu	GGA Gly	ATT Ile	CCA Pro	GGG Gly 40	CCC Pro	ACA Thr	CCT Pro	CTG Leu	CCT Pro 45	TTT Phe	TTG Leu	GGA Gly	144
	AAT Asn	ATT Ile 50	TTG Leu	TCC Ser	TAC Tyr	CAT His	AAG Lys 55	GGC Gly	TTT Phe	TGT Cys	ATG Met	TTT Phe 60	GAC Asp	ATG Met	G <b>AA</b> Glu	TGT Cys	192
50																	

	CAT His 65	AAA Lys	AAG Lys	TAT Tyr	GGA Gly	AAA Lys 70	GTG Val	TGG Trp	GGC Gly	TTT Phe	TAT Tyr 75	GAT Asp	GGT Gly	C <b>AA</b> Gln	CAG Gln	CCT Pro 80	240
5	GTG Val	CTG Leu	GCT Ala	ATC Ile	ACA Thr 85	GAT Asp	CCT Pro	GAC Asp	ATG Met	ATC Ile 90	AAA Lys	ACA Thr	GTG Val	CTA Leu	GTG Val 95	AAA Lys	288
10	GAA Glu	TGT Cys	TAT Tyr	TCT Ser 100	GTC Val	TTC Phe	ACA Thr	AAC Asn	CGG Arg 105	AGG Arg	CCT Pro	TTT Phe	GGT Gly	CCA Pro 110	GTG Val	GGA Gly	336
	TTT Phe	ATG Met	AAA Lys 115	AGT Ser	GCC Ala	ATC Ile	TCT Ser	ATA Ile 120	GCT Ala	GAG Glu	GAT Asp	GAA Glu	GAA Glu 125	TGG Trp	AAG Lys	AGA Arg	384
15	TTA Leu	CGA Arg 130	TCA Ser	TTG Leu	CTG Leu	TCT Ser	CCA Pro 135	ACC Thr	TTC Phe	ACC Thr	AGT Ser	GGA Gly 140	AAA Lys	CTC Leu	AAG Lys	GAG Glu	432
20	ATG Met 145	GTC Val	CCT Pro	ATC Ile	ATT Ile	GCC Ala 150	CAG Gln	TAT Tyr	GGA Gly	GAT Asp	GTG Val 155	TTG Leu	GTG Val	AGA Arg	AAT Asn	CTG Leu 160	480
	AGG Arg	CGG Arg	GAA Glu	GCA Ala	GAG Glu 165	ACA Thr	GGC Gly	AAG Lys	CCT Pro	GTC Val 170	ACC Thr	TTG Leu	ГЛа <b>УУУ</b>	GAC Asp	GTC Val 175	TTT Phe	528
25	GGG Gly	GCC Ala	TAC Tyr	AGC Ser 180	ATG Met	GAT Asp	GTG Val	ATC Ile	ACT Thr 185	AGC Ser	ACA Thr	TCA Ser	TTT Phe	GGA Gly 190	GTG Val	AAC Asn	576
30	ATC Ile	GAC Asp	TCT Ser 195	CTC Leu	AAC Asn	AAT Asn	CCA Pro	CAA Gln 200	GAC Asp	CCC Pro	TTT Phe	GTG Val	GAA Glu 205	AAC Asn	ACC Thr	AAG Lys	624
	<b>AA</b> G Lys	CTT Leu 210	TTA Leu	AGA Arg	TTT Phe	GAT Asp	TTT Phe 215	TTG Leu	GAT Asp	CCA Pro	TTC Phe	TTT Phe 220	CTC Leu	TCA Ser	ATA Ile	ACA Thr	672
35	GTC Val 225	Phe	CCA Pro	TTC Phe	CTC Leu	ATC Ile 230	CCA Pro	ATT Ile	CTT Leu	GAA Glu	GTA Val 235	TTA Leu	AAT Asn	ATC Ile	TGT Cys	GTG Val 240	720
40	TTT Phe	CCA Pro	AGA Arg	GAA Glu	GTT Val 245	ACA Thr	AAT Asn	TTT Phe	TTA Leu	AGA Arg 250	AAA Lys	TCT Ser	GTA Val	AAA Lys	AGG Arg 255	ATG Met	768
	AAA Lys	GAA Glu	AGT Ser	CGC Arg 260	CTC Leu	GAA Glu	GAT Asp	ACA Thr	CAA Gln 265	<b>AA</b> G	CAC His	CGA Arg	GTG Val	GAT Asp 270	TTC Phe	CTT Leu	816
<b>4</b> 5	C <b>A</b> G Gln	CTG Leu	ATG Met 275	Ile	GAC Asp	TCT Ser	CAG Gln	AAT Asn 280	TCA Ser	AAA Lys	G <b>AA</b> Glu	ACT Thr	GAG Glu 285	TCC Ser	CAC His	AAA Lys	864

			GAG Glu						912
5			ACG Thr 310						960
10			GAT Asp						1008
			AAG Lys						1056
15			ATG Met						1104
20			GAG Glu						1152
20			AAA Lys 390						1200
25			AAG Lys						1248
20			AAG Lys						1296
30			GGA Gly						1344
35			CTT Leu						1392
			GAA Glu 470						1440
40			G <b>AA</b> Glu						1488
<b>4</b> 5			GGA Gly	TGA					1512

	(2)	INFO	RMAT	rion	FOR	SEQ	ID 1	10: 8	3:							
5		(	( <i>I</i>	SEQUE A) LE B) TY D) TO	ENGTI PE:	H: 50 amir	)3 am	nino cid		_						
		(ii)	MOI	LECUI	LE TY	PE:	prot	ein								
10		(xi)	SEÇ	QUENC	CE DE	ESCRI	TPTIC	ON: S	SEQ 1	ID NO	D: 8	:				
	Met 1	Ala	Leu	Ile	Pro 5	Asp	Leu	Ala	Met	Glu 10	Thr	Trp	Leu	Leu	Leu 15	Ala
15	Val	Ser	Leu	Val 20	Leu	Leu	Tyr	Leu	Tyr 25	Gly	Thr	His	Ser	His 30	Gly	Leu
	Phe	Lys	Lys 35	Leu	Gly	Ile	Pro	Gly 40	Pro	Thr	Pro	Leu	Pro <b>4</b> 5	Phe	Leu	Gly
20	Asn	Ile 50	Leu	Ser	Tyr	His	Lys 55	Gly	Phe	Cys	Met	Phe 60	Asp	Met	Glu	Cys
	His 65	Lys	Lys	Tyr	Gly	Lys 70	Val	Trp	Gly	Phe	<b>Tyr</b> 75	Asp	Gly	Gln	Gln	Pro 80
25	Val	Leu	Ala	Ile	Thr 85	Asp	Pro	Asp	Met	Ile 90	Lys	Thr	Val	Leu	Val 95	Lys
	Glu	Cys	Tyr	Ser 100	Val	Phe	Thr	Asn	<b>Arg</b> 105	Arg	Pro	Phe	Gly	Pro 110	Val	Gly
30	Phe	Met	Lys 115	Ser	Ala	Ile	Ser	Ile 120	Ala	Glu	Asp	Glu	Glu 125	Trp	Lys	Arg
	Leu	Arg 130	Ser	Leu	Leu	Ser	Pro 135	Thr	Phe	Thr	Ser	Gly 140	Lys	Leu	Lys	Glu
35	Met 145	Val	Pro	Ile	Ile	Ala 150	Gln	Tyr	Gly	Asp	Val 155	Leu	Val	Arg	Asn	Leu 160
	Arg	Arg	Glu	Ala	Glu 165	Thr	Gly	Lys	Pro	<b>V</b> al 170	Thr	Leu	Lys	Asp	Val 175	Phe
40	Gly	Ala	Tyr	Ser 180	Met	Asp	Val	Ile	Thr 185	Ser	Thr	Ser	Phe	Gly 190	Val	Asn
	Ile	Asp	Ser 195	Leu	Asn	Asn	Pro	Gln 200	Asp	Pro	Phe	Val	Glu 205	Asn	Thr	Lys
<b>4</b> 5	Lys	Leu 210	Leu	Arg	Phe	Asp	Phe 215	Leu	Asp	Pro	Phe	Phe 220	Leu	Ser	Ile	Thr
	Val 225	Phe	Pro	Phe	Leu	Ile 230	Pro	Ile	Leu	Glu	Val 235	Leu	Asn	Ile	Cys	Val 240
50	Phe	Pro	Arg	Glu	Val 245	Thr	Asn	Phe	Leu	Arg 250	Lys	Ser	Val	Lys	Arg 255	Met

	Lys	Glu	Ser	Arg 260	Leu	Glu	Asp	Thr	Gln 265	Lys	His	Arg	Val	Asp 270	Phe	Leu
5	Gln	Leu	Met 275	Ile	Asp	Ser	Gln	Asn 280	Ser	Lys	Glu	Thr	Glu 285	Ser	His	Lys
	Ala	Leu 290	Ser	Asp	Leu	Glu	Leu 295	Val	Ala	Gln	Ser	Ile 300	Ile	Phe	Ile	Phe
10	Ala 305	Gly	Tyr	Glu	Thr	Thr 310	Ser	Ser	Val	Leu	Ser 315	Phe	Ile	Met	Tyr	Glu 320
	Leu	Ala	Thr	His	Pro 325	Asp	Val	Gln	Gln	Lys 330	Leu	Gln	Glu	Glu	Ile 335	Asp
15	Ala	Val	Leu	Pro 340	Asn	Lys	Ala	Pro	Pro 345	Thr	Tyr	Asp	Thr	Val 350	Leu	Gln
20	Met	Glu	Tyr 355	Leu	Asp	Met	Val	<b>V</b> al 360	Asn	Glu	Thr	Leu	Arg 365	Leu	Phe	Pro
20	Ile	Ala 370	Met	Arg	Leu	Glu	Arg 375	Val	Cys	Lys	Lys	Asp 380	Val	Glu	Ile	Asn
25	Gly 385	Met	Phe	Ile	Pro	Lys 390	Gly	Trp	Val	Val	<b>M</b> et 395	Ile	Pro	Ser	Tyr	Ala 400
	Leu	His	Arg	Asp	Pro 405	Lys	Tyr	Trp	Thr	Glu 410	Pro	Glu	Lys	Phe	Leu 415	Pro
30	Glu	Arg	Phe	Ser 420	Lys	Lys	Asn	Lys	Asp 425	Asn	Ile	Asp	Pro	Tyr 430	Ile	Tyr
	Thr	Pro	Phe <b>43</b> 5	Gly	Ser	Gly	Pro	Arg 440	Asn	Cys	Ile	Gly	Met 445	Arg	Phe	Ala
35	Leu	Met 450	Asn	Met	Lys	Leu	Ala 455	Leu	Ile	Arg	Val	Leu 460	Gln	Asn	Phe	Ser
	Phe 465	Lys	Pro	Cys	Lys	Glu 470	Thr	Gln	Ile	Pro	Leu 475	Lys	Leu	Ser	Leu	Gly 480
<b>4</b> 0	Gly	Leu	Leu	Gln	Pro 485	Glu	Lys	Pro	Val	Val 490	Leu	Lys	Val	Glu	Ser 495	Arg
	Asp	Gly	Thr	Val 500	Ser	Gly	Ala									
<b>4</b> 5	(2)	INFO	RMAT	TION	FOR	SEQ	ID 1	10: 9	):							
50		(i)	( <i>I</i> (E	QUENC A) LE B) TY C) ST O) TO	ENGTI (PE : TRANI	I: 15 nuc] DEDNE	39 k leic ESS:	ase acio douk	pair 1	îs						

(ix) FEATURE: (A) NAME/KEY: CDS (B) LOCATION: 1..1536 (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 9: ATG CTT TTC CCA ATC TCC ATG TCG GCC ACG GAG TTT CTT CTG GCC TCT 48 Met Leu Phe Pro Ile Ser Met Ser Ala Thr Glu Phe Leu Leu Ala Ser 10 GTC ATC TTC TGT CTG GTA TTC TGG GTA ATC AGG GCC TCA AGA CCT CAG 96 Val Ile Phe Cys Leu Val Phe Trp Val Ile Arg Ala Ser Arg Pro Gln GTC CCC AAA GGC CTG AAG AAT CCA CCA GGG CCA TGG GGC TGG CCT CTG 144 Val Pro Lys Gly Leu Lys Asn Pro Pro Gly Pro Trp Gly Trp Pro Leu 15 ATT GGG CAC ATG CTG ACC CTG GGA AAG AAC CCG CAC CTG GCA CTG TCA 192 Ile Gly His Met Leu Thr Leu Gly Lys Asn Pro His Leu Ala Leu Ser 20 AGG ATG AGC CAG CAG TAT GGG GAC GTG CTG CAG ATC CGA ATT GGC TCC 240 Arg Met Ser Gln Gln Tyr Gly Asp Val Leu Gln Ile Arg Ile Gly Ser ACA CCC GTG GTG GTG CTG AGC GGC CTG GAC ACC ATC CGG CAG GCC CTG 288 Thr Pro Val Val Val Leu Ser Gly Leu Asp Thr Ile Arg Gln Ala Leu 25 GTG CGG CAG GGC GAT GAT TTC AAG GGC CGG CCC GAC CTC TAC ACC TTC 336 Val Arg Gln Gly Asp Asp Phe Lys Gly Arg Pro Asp Leu Tyr Thr Phe 100 105 30 ACC CTC ATC AGT AAT GGT CAG AGC ATG TCC TTC AGC CCA GAC TCT GGA 384 Thr Leu Ile Ser Asn Gly Gln Ser Met Ser Phe Ser Pro Asp Ser Gly 120 CCA GTG TGG GCT GCC CGG CGC CTG GCC CAG AAT GGC CTG AAA AGT Pro Val Trp Ala Ala Arg Arg Leu Ala Gln Asn Gly Leu Lys Ser 35 140 TTC TCC ATT GCC TCT GAC CCA GCC TCC TCA ACC TCC TGC TAC CTG GAA 480 Phe Ser Ile Ala Ser Asp Pro Ala Ser Ser Thr Ser Cys Tyr Leu Glu GAG CAT GTG AGC AAG GAG GCT GAG GTC CTG ATA AGC ACG TTG CAG GAG 528 Glu His Val Ser Lys Glu Ala Glu Val Leu Ile Ser Thr Leu Gln Glu CTG ATG GCA GGG CCT GGG CAC TTT AAC CCC TAC AGG TAT GTG GTA 576 Leu Met Ala Gly Pro Gly His Phe Asn Pro Tyr Arg Tyr Val Val Val 45 185 TCA GTG ACC AAT GTC ATC TGT GCC ATT TGC TTT GGC CGG CGC TAT GAC 624 Ser Val Thr Asn Val Ile Cys Ala Ile Cys Phe Gly Arg Arg Tyr Asp

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				CTT Leu 215						672
5				GGA Gly						720
10				TCC Ser						768
				CAG Gln						816
15				CGG Arg						864
20				GAT Asp 295						912
				GTC Val						960
25				TCC Ser						1008
30				ATC Ile						1056
				CTC Leu						1104
35				ACC Thr 375					TTC Phe	1152
<b>4</b> 0				ACA Thr						1200
				GTC Val						1248
<b>4</b> 5				AAC Asn						1296
50				ATC Ile						1344

	ATC Ile	TTT Phe 450	GGC Gly	ATG Met	GGC Gly	AAG Lys	CGG Arg 455	AAG Lys	TGT Cys	ATC Ile	GGT Gly	GAG Glu 460	ACC Thr	ATT Ile	GCC <b>A</b> la	AGC Ser	1392
5	TGG Trp 465	GAG Glu	GTC Val	TTT Phe	CTC Leu	TTC Phe 470	CTG Leu	GCT Ala	ATC Ile	CTG Leu	CTG Leu 475	CAA Gln	CGG Arg	GTG Val	GAA Glu	TTC Phe 480	1440
10	AGC Ser	GTG Val	CCA Pro	CTG Leu	GGC Gly 485	GTG Val	<b>AA</b> G Lys	GTG Val	GAC Asp	ATG Met 490	ACC Thr	CCC Pro	ATC Ile	TAT Tyr	GGG Gly <b>4</b> 95	CTA Leu	1488
	ACC Thr	ATG Met	AAG Lys	CAT His 500	GCC Ala	TGC Cys	TGT Cys	GAG Glu	CAC His 505	TTC Phe	CAA Gln	ATG Met	CAG Gln	CTG Leu 510	CGC <b>Arg</b>	TCT Ser	1536
15	TAG																1539
	(2)	INFO	OR <b>MA</b> T	rion	FOR	SEQ	ID 1	NO: 3	10:								
20			( <i>I</i>	SEQUI A) LI B) T' O) T(	ENGTI PE:	H: 53 amir	L2 ar	mino cid									
				LECUI			-										
25	Mot										0: 10 Glu		I.eu	Len	Ala	Ser	
25	1	Leu	Phe	Pro	Ile 5	Ser	Met	Ser	Ala	Thr 10	Glu	Phe			15		
25	1		Phe	Pro	Ile 5	Ser	Met	Ser	Ala	Thr 10	Glu	Phe			15		
25 30	1 Val Val	Leu Ile Pro	Phe Phe Lys	Pro Cys 20 Gly	Ile 5 Leu Leu	Ser Val Lys	Met Phe Asn	Ser Trp Pro 40	Ala Val 25 Pro	Thr 10 Ile Gly	Glu Arg Pro	Phe Ala Trp	Ser Gly 45	Arg 30 Trp	Pro	Gln Leu	
	1 Val Val	Leu Ile	Phe Phe Lys	Pro Cys 20 Gly	Ile 5 Leu Leu	Ser Val Lys	Met Phe Asn	Ser Trp Pro 40	Ala Val 25 Pro	Thr 10 Ile Gly	Glu Arg Pro	Phe Ala Trp	Ser Gly 45	Arg 30 Trp	Pro	Gln Leu	
	1 Val Val	Leu Ile Pro Gly	Phe Phe Lys 35	Pro Cys 20 Gly Met	Ile 5 Leu Leu	Ser Val Lys Thr	Met Phe Asn Leu 55	Ser Trp Pro 40 Gly	Val 25 Pro	Thr 10 Ile Gly Asn	Glu Arg Pro	Phe Ala Trp His 60	Ser Gly 45 Leu	Arg 30 Trp Ala	Pro Pro Leu	Gln Leu Ser	
30	Val Val Ile Arg 65	Leu Ile Pro Gly	Phe Phe Lys 35 His	Pro Cys 20 Gly Met	Ile 5 Leu Leu Gln	Ser  Val  Lys  Thr  Tyr 70	Met Phe Asn Leu 55	Ser Trp Pro 40 Gly Asp	Ala Val 25 Pro Lys Val	Thr 10 Ile Gly Asn Leu	Glu Arg Pro Pro Gln 75	Phe Ala Trp His 60	Ser Gly 45 Leu Arg	Arg 30 Trp Ala	Pro Pro Leu Gly	Gln Leu Ser Ser 80	
30	Val Val Ile Arg 65	Leu Ile Pro Gly 50 Met	Phe Lys 35 His Ser Val	Pro Cys 20 Gly Met Gln Val	Ile 5 Leu Leu Gln Val	Ser Val Lys Thr Tyr 70 Leu	Met Phe Asn Leu 55 Gly Ser	Ser Trp Pro 40 Gly Asp	Ala Val 25 Pro Lys Val Leu	Thr 10 Ile Gly Asn Leu	Glu Arg Pro Pro Gln 75 Thr	Phe Ala Trp His 60 Ile	Ser Gly 45 Leu Arg	Arg 30 Trp Ala Ile	Pro Pro Leu Gly Ala 95	Gln Leu Ser Ser 80 Leu	
30 35	Val Val Ile Arg 65 Thr	Leu Ile Pro Gly 50 Met	Phe Lys 35 His Ser Val	Pro Cys 20 Gly Met Gln Val Gly 100	Ile 5 Leu Leu Gln Val 85 Asp	Ser Val Lys Thr Tyr 70 Leu Asp	Met Phe Asn Leu 55 Gly Ser Phe	Ser Trp Pro 40 Gly Asp Gly Lys	Val 25 Pro Lys Val Leu Gly 105	Thr 10 Ile Gly Asn Leu Asp 90 Arg	Glu Arg Pro Pro Gln 75 Thr	Phe Ala Trp His 60 Ile Ile Asp	Ser Gly 45 Leu Arg Arg	Arg 30 Trp Ala Ile Gln Tyr 110	Pro Pro Leu Gly Ala 95 Thr	Gln Leu Ser Ser 80 Leu Phe	

	Phe 145	Ser	Ile	Ala	Ser	Asp 150	Pro	Ala	Ser	Ser	Thr 155	Ser	Cys	Tyr	Leu	Glu 160
5	Glu	His	Val	Ser	Lys 165	Glu	Ala	Glu	Val	Leu 170	Ile	Ser	Thr	Leu	Gln 175	Glu
	Leu	Met	Ala	Gly 180	Pro	Gly	His	Phe	Asn 185	Pro	Tyr	Arg	Tyr	Val 190	Val	Val
10	Ser	Val	Thr 195	Asn	Val	lle	Cys	Ala 200	Ile	Cys	Phe	Gly	<b>A</b> rg 205	Arg	Tyr	Asp
	His	Asn 210	His	Gln	Glu	Leu	Leu 215	Ser	Leu	Val	Asn	Leu 220	Asn	Asn	Asn	Phe
15	Gly 225	Glu	Val	Val	Gly	Ser 230	Gly	Asn	Pro	Ala	Asp 235	Phe	Ile	Pro	Ile	Leu 2 <b>4</b> 0
	Arg	Tyr	Leu	Pro	Asn 245	Pro	Ser	Leu	Asn	Ala 250	Phe	ГÀЗ	Asp	Leu	Asn 255	Glu
20	Lys	Phe	Tyr	Ser 260	Phe	Met	Gln	Lys	Met 265	Val	Lys	Glu	His	Tyr 270	Lys	Thr
	Phe	Glu	Lys 275	Gly	His	Ile	Arg	Asp 280	Ile	Thr	Asp	Ser	Leu 285	Ile	Glu	His
25	Cys	Gln 290	Glu	Lys	Gln	Leu	Asp 295	Glu	Asn	Ala	Asn	Val 300	Gln	Leu	Ser	Asp
	Glu 305	Lys	Ile	Ile	Asn	Ile 310	Val	Leu	Asp	Leu	Phe 315	Gly	Ala	Gly	Phe	Asp 320
30	Thr	Val	Thr	Thr	Ala 325	Ile	Ser	Trp	Ser	Leu 330	Met	Tyr	Leu	Val	Met 335	Asn
	Pro	Arg	Val	Gln 340	Arg	Lys	Ile	Gln	Glu 3 <b>4</b> 5	Glu	Leu	Asp	Thr	Val 350	Ile	Gly
35	Arg	Ser	Arg 355	Arg	Pro	Arg	Leu	Ser 360	Asp	Arg	Ser	His	Leu 365	Pro	Туr	Met ,
	Glu	Ala 370	Phe	Ile	Leu	Glu	Thr 375	Phe	Arg	His	Ser	Ser 380	Phe	Val	Pro	Phe
40	Thr 385	Ile	Pro	His	Ser	Thr 390	Thr	Arg	Asp	Thr	Ser 395	Leu	Lys	Gly	Phe	Tyr 400
	Ile	Pro	Lys	Gly	Arg 405	Cys	Val	Phe	Val	Asn 410	Gln	Trp	Gln	Ile	<b>As</b> n <b>41</b> 5	His
<b>4</b> 5	Asp	Gln	Lys	Leu 420	Trp	Val	Asn	Pro	Ser 425	Glu	Phe	Leu	Pro	Glu 430	Arg	Phe
	Leu	Thr	Pro 435	Asp	Gly	Ala	Ile	Asp 440	Lys	Val	Leu	Ser	Glu 445	Lys	Val	Ile
50	Ile	Phe 450	Gly	Met	Gly	Lys	Arg 455	Lys	Cys	Ile	Gly	Glu 460	Thr	Ile	Ala	Ser

	Trp 465	Glu	Val	Phe	Leu	Phe 470	Leu	Ala	Ile	Leu	Leu <b>4</b> 75	Gln	Arg	Val	Glu	Phe 480	
5	Ser	Val	Pro	Leu	Gly 485	Val	Lys	Val	Asp	Met 490	Thr	Pro	Ile	Tyr	Gly 495	Leu	
	Thr	Met	Lys	His 500	Ala	Cys	Cys	Glu	His 505	Phe	Gln	Met	Gln	Leu 510	Arg	Ser	
10	(2)		ORMAT														
15		(i)	(E		ENGTI (PE : TRANI	H: 15 nucl DEDNE	39 l leic ESS:	acio doub	pai:	cs							
20		(ix)		ATURI A) NA B) LA	ME/I			1536									
		(xi)	SEÇ	QUENC	CE DI	ESCR	PTI	ON: S	SEQ :	D NO	): 1	l:					
25	ATG Met	CTT Leu	TTC Phe	CCA Pro	ATC Ile 5	TCC Ser	ATG Met	TCG Ser	GCC Ala	ACG Thr 10	GAG Glu	TTT Phe	CTT Leu	CTG Leu	GCC Ala 15	TCT Ser	<b>4</b> 8
	GTC Val	ATC Ile	TTC Phe	TGT Cys 20	CTG Leu	GTA Val	TTC Phe	TGG Trp	GTA Val 25	ATC Ile	AGG Arg	GCC Ala	TCA Ser	AGA Arg 30	CCT Pro	CAG Gln	96
30	GTC Val	CCC Pro	AAA Lys 35	GGC Gly	CTG Leu	AAG Lys	AAT Asn	CCA Pro 40	CCA Pro	GGG Gly	CCA Pro	TGG Trp	GGC Gly 45	TGG Trp	CCT Pro	CTG Leu	144
<b>3</b> 5	ATT Ile	GGG Gly 50	CAC His	ATG Met	CTG Leu	ACC Thr	CTG Leu 55	GGA Gly	AAG Lys	AAC Asn	CCG Pro	CAC His 60	CTG Leu	GCA Ala	CTG Leu	TCA Ser	192
	AGG Arg 65	ATG Met	AGC Ser	CAG Gln	CAG Gln	TAT Tyr 70	GGG Gly	GAC Asp	GTG Val	CTG Leu	CAG Gln 75	ATC Ile	CGA Arg	ATT Ile	GGC Gly	TCC Ser 80	240
40	ACA Thr	CCC Pro	GTG Val	GTG Val	GTG Val 85	CTG Leu	AGC Ser	GGC Gly	CTG Leu	GAC Asp 90	ACC Thr	ATC Ile	CGG Arg	CAG Gln	GCC Ala 95	CTG Leu	288
	GTG Val	CGG <b>A</b> rg	CAG Gln	GGC Gly 100	GAT Asp	GAT <b>A</b> sp	TTC Phe	AAG Lys	GGC Gly 105	CGG <b>A</b> rg	CCC Pro	GAC Asp	CTC Leu	TAC Tyr 110	ACC Thr	TTC Phe	336
<b>4</b> 5	ACC Thr	CTC Leu	ATC Ile 115	AGT Ser	AAT Asn	GGT Gly	CAG Gln	AGC Ser 120	ATG Met	TCC Ser	TTC Phe	AGC Ser	CCA Pro 125	GAC Asp	TCT Ser	GGA Gly	384
50																	

0

	CCA Pro	GTG Val 130	TGG Trp	GCT Ala	GCC Ala	CGC Arg	CGG Arg 135	CGC <b>A</b> rg	CTG Leu	GCC Ala	CAG Gln	AAT Asn 140	GGC Gly	CTG Leu	AAA Lys	AGT Ser	432
5	TTC Phe 145	TCC Ser	ATT Ile	GCC Ala	TCT Ser	GAC Asp 150	CCA Pro	GCC Ala	TCC Ser	TCA Ser	ACC Thr 155	TCC Ser	TGC Cys	TAC Tyr	CTG Leu	GAA Glu 160	480
10	GAG Glu	CAT His	GTG Val	AGC Ser	AAG Lys 165	GAG Glu	GCT Ala	GAG Glu	GTC Val	CTG Leu 170	ATA Ile	AGC Ser	ACG Thr	TTG Leu	CAG Gln 175	GAG Glu	528
	CTG Leu	ATG Met	GCA Ala	GGG Gly 180	CCT Pro	GGG Gly	CAC His	TTT Phe	AAC Asn 185	CCC Pro	TAC Tyr	AGG Arg	TAT Tyr	GTG Val 190	GTG Val	GTA Val	576
15	TCA Ser	GTG Val	ACC Thr 195	AAT Asn	GTC Val	ATC Ile	TGT Cys	GCC Ala 200	ATT Ile	TGC Cys	TTT Phe	GGC Gly	CGG Arg 205	CGC <b>A</b> rg	TAT Tyr	GAC Asp	624
20					GAA Glu												672
	GGG Gly 225	GAG Glu	GTG Val	GTT Val	GGC Gly	TCT Ser 230	GGA Gly	AAC Asn	CCA Pro	GCT Ala	GAC Asp 235	TTC Phe	ATC Ile	CCT Pro	ATT Ile	CTT Leu 240	720
25	CGC Arg	TAC Tyr	CTA Leu	CCC Pro	AAC Asn 245	CCT Pro	TCC Ser	CTG Leu	AAT Asn	GCC Ala 250	TTC Phe	AAG Lys	GAC Asp	CTG Leu	AAT Asn 255	GAG Glu	768
30					TTC Phe												816
	TTT Phe	GAG Glu	AAG Lys 275	GGC Gly	CAC His	ATC Ile	CGG Arg	GAC Asp 280	ATC Ile	ACA Thr	GAC Asp	AGC Ser	CTG Leu 285	ATT Ile	GAG Glu	CAC His	864
35					CAG Gln							_	_			GAT Asp	912
40	GAG Glu 305	AAG Lys	ATC Ile	ATT Ile	AAC Asn	ATC Ile 310	GTC Val	TTG Leu	GAC Asp	CTC Leu	TTT Phe 315	GGA Gly	GCT Ala	GGG Gly	TTT Phe	GAC Asp 320	960
				Thr	GCT Ala 325	Ile	Ser	$\mathtt{Trp}$	Ser	Leu	Met	Tyr	Leu	Val		Asn	1008
<b>4</b> 5	CCC Pro	AGG Arg	GTA Val	CAG Gln 340	AGA Arg	rya YyG	ATC Ile	CAA Gln	GAG Glu 345	GAG Glu	CTC Leu	GAC Asp	ACA Thr	GTG Val 350	ATT	GGC Gly	1056
50					CCC Pro												1104

	GAG Glu	GCC Ala 370	TTC Phe	ATC Ile	CTG Leu	G <b>A</b> G Glu	ACC Thr 375	TTC Phe	CGA Arg	CAC His	TCT Ser	TCC Ser 380	TTC Phe	GTC Val	CCC Pro	TTC Phe	1152
5	ACC Thr 385	ATC Ile	CCC Pro	CAC His	AGC Ser	ACA Thr 390	ACA Thr	AGA Arg	GAC Asp	ACA Thr	AGT Ser 395	TTG Leu	AAA Lys	GGC Gly	TTT Phe	TAC Tyr 400	1200
10	ATC Ile	CCC Pro	AAG Lys	GGG Gly	CGT Arg 405	TGT Cys	GTC Val	TTT Phe	GTA Val	AAC Asn 410	CAG Gln	TGG Trp	CAG Gln	ATC Ile	AAC Asn 415	CAT His	1248
	GAC Asp	CAG Gln	AAG Lys	CTA Leu 420	TGG Trp	GTC Val	AAC Asn	CCA Pro	TCT Ser 425	GAG Glu	TTC Phe	CTA Leu	CCT Pro	GAA Glu 430	CGG Arg	TTT Phe	1296
15	CTC Leu	ACC Thr	CCT Pro 435	GAT Asp	GGT Gly	GCT Ala	ATC Ile	GAC Asp 440	AAG Lys	GTG Val	TTA Leu	AGT Ser	GAG Glu 445	AAG Lys	GTG Val	ATT Ile	1344
20	ATC Ile	TTT Phe 450	GGC Gly	ATG Met	GGC Gly	AAG Lys	CGG Arg 455	AAG Lys	TGT Cys	ATC Ile	GGT Gly	GAG Glu 460	ACC Thr	ATT Ile	GCC <b>Al</b> a	CGC <b>A</b> rg	1392
	TGG Trp 465	GAG Glu	GTC Val	TTT Phe	CTC Leu	TTC Phe 470	CTG Leu	GCT Ala	ATC Ile	CTG Leu	CTG Leu 475	CAA Gln	CGG Arg	GTG Val	GAA Glu	TTC Phe 480	1440
25	AGC Ser	GTG Val	CCA Pro	CTG Leu	GGC Gly 485	GTG Val	AAG Lys	GTG Val	GAC Asp	ATG Met 490	ACC Thr	CCC Pro	ATC Ile	TAT Tyr	GGG Gly 495	CTA Leu	1488
30	ACC Thr	ATG Met	AAG Lys	CAT His 500	GCC Ala	TGC Cys	TGT Cys	GAG Glu	CAC His 505	TTC Phe	CAA Gln	ATG Met	CAG Gln	CTG Leu 510	CGC <b>A</b> rg	TCT Ser	1536
	TAG																1539
	(2)	INFO	RMAT	rion	FOR	SEQ	ID I	<b>1</b> 0: 1	L2:								
35		(	( <i>I</i>	SEQUE A) LE B) TY	ENGTI PE :	H: 52 amir	L2 ar	nino cid									

- (ii) MOLECULE TYPE: protein
  - (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 12:

Met Leu Phe Pro Ile Ser Met Ser Ala Thr Glu Phe Leu Leu Ala Ser 1 5 10 15

Val Ile Phe Cys Leu Val Phe Trp Val Ile Arg Ala Ser Arg Pro Gln 20 25 30

Val Pro Lys Gly Leu Lys Asn Pro Pro Gly Pro Trp Gly Trp Pro Leu 35 40 45

50

	Ile	Gly 50	His	Met	Leu	Thr	Leu 55	Gly	Lys	Asn	Pro	His 60	Leu	Ala	Leu	Ser
5	Arg 65	Met	Ser	Gln	Gln	Tyr 70	Gly	Asp	Val	Leu	Gln 75	Ile	Arg	Ile	Gly	Ser 80
	Thr	Pro	Val	Val	Val 85	Leu	Ser	Gly	Leu	Asp 90	Thr	Ile	Arg	Gln	Ala 95	Leu
10	Val	Arg	Gln	Gly 100	Asp	Asp	Phe	Lys	Gly 105	Arg	Pro	Asp	Leu	Tyr 110	Thr	Phe
	Thr	Leu	Ile 115	Ser	Asn	Gly	Gln	Ser 120	Met	Ser	Phe	Ser	Pro 125	Asp	Ser	Gly
15	Pro	Val 130	Trp	Ala	Ala	Arg	Arg 135	Arg	Leu	Ala	Gln	Asn 140	Gly	Leu	Lys	Ser
20	Phe 145	Ser	Ile	Ala	Ser	Asp 150	Pro	Ala	Ser	Ser	Thr 155	Ser	Суѕ	Tyr	Leu	Glu 160
20	Glu	His	Val	Ser	Lys 165	Glu	Ala	Glu	Val	Leu 170	Ile	Ser	Thr	Leu	Gln 175	Glu
25	Leu	Met	Ala	Gly 180	Pro	Gly	His	Phe	Asn 185	Pro	Tyr	Arg	Tyr	Val 190	Val	Val
	Ser	Val	Thr 195	Asn	Val	Ile	Cys	Ala 200	Ile	Суѕ	Phe	Gly	Arg 205	Arg	Tyr	Asp
30	His	Asn 210	His	Gln	Glu	Leu	Leu 215	Ser	Leu	Val	Asn	Leu 220	Asn	Asn	Asn	Phe
	Gly 225	Glu	Val	Val	Gly	Ser 230	Gly	Asn	Pro	Ala	Asp 235	Phe	Ile	Pro	Ile	Leu 240
35	Arg	Tyr	Leu	Pro	Asn 245	Pro	Ser	Leu	Asn	Ala 250	Phe	Lys	Asp	Leu	Asn 255	Glu
	Lys	Phe	Tyr	Ser 260	Phe	Met	Gln	Lys	Met 265	Val	Lys	Glu	His	Tyr 270	Lys	Thr
40	Phe	Glu	Lys 275	Gly	His	Ile	Arg	Asp 280	Ile	Thr	Asp	Ser	Leu 285	Ile	Glu	His
	Cys	Gln 290	Glu	Lys	Gln	Leu	Asp 295	Glu	Asn	Ala	Asn	Val 300	Gln	Leu	Ser	Asp
<b>4</b> 5	Glu 305	Lys	Ile	Ile	Asn	Ile 310	Val	Leu	Asp	Leu	Phe 315	Gly	Ala	Gly	Phe	Asp 320
	Thr	Val	Thr	Thr	Ala 325	Ile	Ser	Trp	Ser	Leu 330	Met	Tyr	Leu	Val	Met 335	Asn
50	Pro	Arg	Val	Gln 340	Arg	Lys	Ile	Gln	Glu 3 <b>4</b> 5	Glu	Leu	Asp	Thr	<b>V</b> al	Ile	Gly

	Arg	Ser	Arg 355	Arg	Pro	Arg	Leu	Ser 360	Asp	Arg	Ser	His	Leu 365	Pro	Tyr	Met		
5	Glu	Ala 370	Phe	Ile	Leu	Glu	Thr 375	Phe	Arg	His	Ser	Ser 380	Phe	Val	Pro	Phe		
	Thr 385	Ile	Pro	His	Ser	Thr 390	Thr	Arg	Asp	Thr	Ser 395	Leu	Lys	Gly	Phe	Tyr <b>4</b> 00		
10	Ile	Pro	Lys	Gly	Arg 405	Cys	Val	Phe	Val	Asn 410	Gln	Trp	Gln	Ile	Asn 415	His		
	Asp	Gln	Lys	Leu 420	Trp	Val	Asn	Pro	Ser 425	Glu	Phe	Leu	Pro	Glu 430	Arg	Phe		
15	Leu	Thr	Pro 435	Asp	Gly	Ala	Ile	Asp 440	Lys	Val	Leu	Ser	Glu <b>44</b> 5	Lys	Val	Ile		
	Ile	Phe 450	Gly	Met	Gly	Lys	<b>A</b> rg <b>4</b> 55	Lys	Cys	Ile	Gly	Glu 460	Thr	Ile	Ala	Arg		
20	Trp	Glu	Val	Phe	Leu	Phe 470	Leu	Ala	Ile	Leu	Leu 475	Gln	Arg	Val	Glu	Phe 480		
	Ser	Val	Pro	Leu	Gly 485	Val	Lys	Val	Asp	Met 490	Thr	Pro	Ile	Tyr	Gly 495	Leu		
25	Thr	Met	Lys	His 500	Ala	Cys	Cys	Glu	His 505	Phe	Gln	Met	Gln	Leu 510	Arg	Ser		
	(2)	INFO	RMAT	NOI	FOR	SEQ	ID N	<b>1</b> 0: 1	13:									
30		(i)	( <i>I</i> (E	QUENCA) LE B) TY C) ST D) TO	ENGTI (PE : [RANI	H: 15 nucl DED <b>N</b> E	39 k leic ESS:	ase acio doul	pai:	cs								
<b>3</b> 5		(ix)	( ]	ATURE A) NA B) LO	AME/I			1536										
				QUENC												_		
40	ATG Met 1	CTT Leu	TTC Phe	CCA Pro	ATC Ile 5	TCC Ser	ATG Met	TCG Ser	GCC Ala	ACG Thr 10	GAG Glu	TTT	CTT Leu	CTG Leu	GCC Ala 15	TCT Ser		48
<b>4</b> 5	GTC Val	ATC Ile	TTC Phe	TGT Cys 20	CTG Leu	GTA Val	TTC Phe	TGG Trp	GTA Val 25	ATC Ile	AGG Arg	GCC Ala	TCA Ser	AGA Arg 30	CCT Pro	CAG Gln		96
		CCC Pro															Ē	144
50																		

				CTG Leu 55	_				_			192
5				GGG Gly								240
10				AGC Ser								288
				TTC Phe								336
15				CAG Gln			_					384
20				CGG Arg 135				_				432
				CCA Pro								480
25				GCT Ala								528
30				CAC His								576
				TGT Cys								624
35				CTT Leu 215							TTC ' Phe	672
<b>4</b> 0				GGA Gly				_		_		720
				TCC Ser								768
<b>4</b> 5				CAG Gln								816
				CGG Arg								864

	TGT Cys	CAG Gln 290	GAG Glu	<b>AA</b> G Lys	CAG Gln	CTG Leu	GAT Asp 295	GAG Glu	AAC Asn	GCC Ala	AAT Asn	GTC Val 300	CAG Gln	CTG Leu	TCA Ser	GAT Asp	912
5	GAG Glu 305	AAG Lys	ATC Ile	ATT Ile	AAC Asn	ATC Ile 310	GTC Val	TTG Leu	GAC Asp	CTC Leu	TTT Phe 315	GGA Gly	GCT Ala	GGG Gly	TTT Phe	GAC Asp 320	960
10	ACA Thr	GTC Val	ACA Thr	ACT Thr	GCT Ala 325	ATC Ile	TCC Ser	TGG Trp	AGC Ser	CTC Leu 330	ATG Met	TAT Tyr	TTG Leu	GTG Val	ATG Met 335	AAC Asn	1008
	CCC Pro	AGG Arg	GTA Val	CAG Gln 340	AGA Arg	<b>AA</b> G Lys	ATC Ile	CAA Gln	GAG Glu 345	GAG Glu	CTC Leu	GAC Asp	ACA Thr	GTG Val 350	ATT Ile	GGC Gly	1056
15	AGG Arg	TCA Ser	CGG Arg 355	CGG Arg	CCC Pro	CGG <b>A</b> rg	CTC Leu	TCT Ser 360	GAC Asp	AGA Arg	TCC Ser	CAT His	CTG Leu 365	CCC Pro	TAT Tyr	ATG Met	1104
20	G <b>A</b> G Glu	GCC Ala 370	TTC Phe	ATC Ile	CTG Leu	GAG Glu	ACC Thr 375	TTC Phe	CGA Arg	CAC His	TCT Ser	TCC Ser 380	TTC Phe	GTC Val	CCC Pro	TTC Phe	1152
	ACC Thr 385	ATC Ile	CCC Pro	CAC His	AGC Ser	ACA Thr 390	ACA Thr	AGA Arg	GAC Asp	ACA Thr	AGT Ser 395	TTG Leu	AAA Lys	GGC Gly	TTT Phe	TAC Tyr 400	1200
25	ATC Ile	CCC Pro	AAG Lys	GGG Gly	CGT Arg 405	TGT Cys	GTC Val	TTT Phe	GTA Val	AAC Asn 410	CAG Gln	TGG Trp	CAG Gln	ATC Ile	AAC Asn 415	CAT His	1248
30	GAC Asp	CAG Gln	AAG Lys	CTA Leu 420	TGG Trp	GTC Val	AAC Asn	CCA Pro	TCT Ser 425	GAG Glu	TTC Phe	CTA Leu	CCT Pro	GAA Glu 430	CGG <b>A</b> rg	TTT Phe	1296
	CTC Leu	ACC Thr	CCT Pro 435	GAT Asp	GGT Gly	GCT Ala	ATC Ile	GAC Asp 440	AAG Lys	GTG Val	TTA Leu	AGT Ser	GAG Glu 445	AAG Lys	GTG Val	ATT Ile	1344
35	ATC Ile	TTT Phe 450	GGC Gly	ATG Met	GGC Gly	AAG Lys	CGG Arg 455	AAG Lys	TGT Cys	ATC Ile	GGT Gly	GAG Glu 460	ACC Thr	GTT Val	GCC Ala	CGC <b>A</b> rg	1392
<b>4</b> 0	TGG Trp 465	GAG Glu	GTC Val	TTT Phe	CTC Leu	TTC Phe 470	CTG Leu	GCT Ala	ATC Ile	CTG Leu	CTG Leu 475	CAA Gln	CGG <b>Ar</b> g	GTG Val	GAA Glu	TTC Phe 480	1440
	AGC Ser	GTG Val	CCA Pro	CTG Leu	GGC Gly 485	GTG Val	AAG Lys	GTG Val	GAC Asp	ATG Met 490	ACC Thr	CCC Pro	ATC Ile	TAT Tyr	GGG Gly 495	CTA Leu	1488
<b>4</b> 5									CAC His 505								1536
	TAG																1539

	(2)	INT	ORMA	11014	FOR	SEQ	10	NO:	14:							
5			(	A) L B) T	ENCE ENGT YPE : OPOL	H: 5 ami	12 a no a	mino cid		_						
		(ii	) MO	LECU	LE T	YPE:	pro	tein								
		(xi	) SE	QUEN	CE D	ESCR	IPTI	ON:	SEQ	ID N	0: 1	4:				
10	Met 1	Leu	Phe	Pro	Ile 5	Ser	Met	Ser	Ala	Thr 10	Glu	Phe	Leu	Leu	Ala 15	
_	Val	Ile	Phe	Cys 20	Leu	Val	Phe	Trp	Val 25	Ile	Arg	Ala	Ser	Arg 30	Pro	Glr
15	Val	Pro	Lys 35	Gly	Leu	Lys	Asn	Pro 40	Pro	Gly	Pro	Trp	Gly 45	Trp	Pro	Leu
	Ile	Gly 50	His	Met	Leu	Thr	Leu 55	Gly	Lys	<b>A</b> sn	Pro	His 60	Leu	Ala	Leu	Ser
20	Arg 65	Met	Ser	Gln	Gln	Туг 70	Gly	Asp	Val	Leu	Gln 75	Ile	Arg	Ile	Gly	Ser 80
	Thr	Pro	Val	Val	Val 85	Leu	Ser	Gly	Leu	Asp 90	Thr	Ile	Arg	Gln	Ala 95	Leu
25	Val	Arg	Gln	Gly 100	Asp	Asp	Phe	Lys	Gly 105	Arg	Pro	Asp	Leu	Tyr 110	Thr	Phe
	Thr	Leu	Ile 115	Ser	Asn	Gly	Gln	Ser 120	Met	Ser	Phe	Ser	Pro 125	Asp	Ser	Gly
30	Pro	Val 130	Trp	Ala	Ala	Arg	Arg 135	Arg	Leu	Ala	Gln	Asn 140	Gly	Leu	Lys	Ser
	Phe 1 <b>4</b> 5	Ser	Ile	Ala	Ser	Asp 150	Pro	Ala	Ser	Ser	Thr 155	Ser	Cys	Tyr	Leu	Glu 160
35	Glu	His	Val	Ser	Lув 165	Glu	Ala	Glu	Val	Leu 170	Ile	Ser	Thr	Leu	Gln 175	Glu
	Leu	Met	Ala	Gly 180	Pro	Gly	His	Phe	Asn 185	Pro	Tyr	Arg	Tyr	<b>Val</b> 190	Val	Val
40	Ser	Val	Thr 195	Asn	Val	Ile	Суѕ	Ala 200	Ile	Cys	Phe	Gly	Arg 205	Arg	Tyr	Asp
	His	Asn 210	His	Gln	Glu	Leu	Leu 215	Ser	Leu	Val	Asn	Leu 220	Asn	Asn	Asn	Phe
<b>4</b> 5	Gly 225	Glu	Val	Val	Gly	Ser 230	Gly	Asn	Pro	Ala	Asp 235	Phe	Ile	Pro	Ile	Leu 240
	Arg	Tyr	Leu	Pro	Asn 245	Pro	Ser	Leu	Asn	Ala 250	Phe	Lys	Asp	Leu	Asn 255	Glu
50																

	Lys	Phe	Tyr	Ser 260	Phe	Met	Gln	Lys	Met 265	Val	Lys	Glu	His	Tyr 270	Lys	Thr
5	Phe	Glu	Lys 275	Gly	His	Ile	Arg	Asp 280	Ile	Thr	Asp	Ser	Leu 285	Ile	Glu	His
	Cys	Gln 290	Glu	Lys	Gln	Leu	Asp 295	Glu	Asn	Ala	Asn	Val 300	Gln	Leu	Ser	Asp
10	Glu 305	Lys	Ile	Ile	Asn	Ile 310	Val	Leu	Asp	Leu	Phe 315	Gly	Ala	Gly	Phe	Asp 320
	Thr	Val	Thr	Thr	Ala 325	Ile	Ser	Trp	Ser	Leu 330	Met	Tyr	Leu	Val	Met 335	Asn
15	Pro	Arg	Val	Gln 340	Arg	Lys	Ile	Gln	Glu 345	Glu	Leu	Asp	Thr	<b>Val</b> 350	Ile	Gly
	Arg	Ser	Arg 355	Arg	Pro	Arg	Leu	Ser 360	Asp	Arg	Ser	His	Leu 365	Pro	Tyr	Met
20	Glu	Ala 370	Phe	Ile	Leu	Glu	Thr 375	Phe	Arg	His	Ser	Ser 380	Phe	Val	Pro	Phe
<b>~</b> r	Thr 385	Ile	Pro	His	Ser	Thr 390	Thr	Arg	Asp	Thr	Ser 395	Leu	Lys	Gly	Phe	Tyr 400
25	Ile	Pro	Lys	Gly	Arg 405	Cys	Val	Phe	Val	Asn 410	Gln	Trp	Gln	Ile	Asn 415	His
30	Asp	Gln	Lys	Leu 420	Trp	Val	Asn	Pro	Ser 425	Glu	Phe	Leu	Pro	Glu 430	Arg	Phe
	Leu	Thr	Pro 435	Asp	Gly	Ala	Ile	Asp 440	Lys	Val	Leu	Ser	Glu 445	Lys	Val	Ile
35	Ile	Phe 450	Gly	Met	Gly	Lys	Arg 455	Lys	Cys	Ile	Gly	Glu 460	Thr	Val	Ala	Arg
	Trp 465	Glu	Val	Phe	Leu	Phe 470	Leu	Ala	Ile	Leu	Leu 475	Gln	Arg	Val	Glu	Phe 480
<b>4</b> 0	Ser	Val	Pro	Leu	Gly 485	Val	Lys	Val	Asp	Met 490	Thr	Pro	Ile	Tyr	Gly 495	Leu
	Thr	Met	Lys	His 500	Ala	Cys	Cys	Glu	His 505	Phe	Gln	Met	Gln	Leu 510	Arg	Ser
<b>4</b> 5	(2)	INF	ORMA'	rion	FOR	SEQ	ID I	<b>10:</b> 3	15:							

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 1485 base pairs

(B) TYPE: nucleic acid

(C) STRANDEDNESS: double

(D) TOPOLOGY: linear

55

(ix) FEATURE:

(A) NAME/KEY: CDS
(B) LOCATION: 1..1482

5		(xi	) SE	OUEN	CE DI	ESCR	IP <b>T</b> I	ON: 5	SEQ .	ID N	D: 1	5:					
	ATG Met	CTG Leu	GCC Ala	TCA Ser	GGG Gly 5	ATG Met	CTT Leu	CTG Leu	GTG Val	GCC Ala 10	TTG Leu	CTG Leu	GTC Val	TGC Cys	CTG Leu 15	ACT Thr	48
10							GTT Val										96
15							CCA Pro										144
	CTG Leu	AAC Asn 50	ACA Thr	GAG Glu	CAG Gln	ATG Met	TAC Tyr 55	AAC Asn	TCC Ser	CTC <b>L</b> eu	ATG Met	AAG Lys 60	ATC Ile	AGT Ser	GAG Glu	CGC <b>A</b> rg	192
20							ATT Ile										240
25							GTC Val										288
							GAG Glu										336
30							AGC Ser										384
35							CTG Leu 135										432
							GAG Glu										480
40	CGG Arg	GGC Gly	ACT Thr	GGC Gly	GGC Gly 165	GCC Ala	AAT Asn	ATC Ile	GAT Asp	CCC Pro 170	ACC Thr	TTC Phe	TTC Phe	CTG Leu	AGC Ser 175	CGC <b>A</b> rg	528
<b>4</b> 5							AGC Ser										576
							CTG Leu										624

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	TTC Phe	CAG Gln 210	TTC Phe	ACG Thr	TCA Ser	ACC Thr	TCC Ser 215	ACG Thr	GGG Gly	CAG Gln	CTC Leu	TAT Tyr 220	GAG Glu	ATG Met	TTC Phe	TCT Ser	672
5	TCG Ser 225	GTG Val	ATG Met	AAA Lys	CAC His	CTG Leu 230	CCA Pro	GGA Gly	CCA Pro	CAG Gln	CAA Gln 235	CAG Gln	GCC Ala	TTT Phe	CAG Gln	TTG Leu 240	720
10	CTG Leu	CAA Gln	GGG Gly	CTG Leu	GAG Glu 245	GAC Asp	TTC Phe	ATA Ile	GCC Ala	AAG Lys 250	AAG Lys	GTG Val	GAG Glu	CAC His	AAC Asn 255	CAG Gln	768
	CGC Arg	ACG Thr	CTG Leu	GAT Asp 260	CCC Pro	AAT Asn	TCC Ser	CCA Pro	CGG Arg 265	GAC Asp	TTC Phe	ATT Ile	GAC Asp	TCC Ser 270	TTT Phe	CTC Leu	816
15	ATC Ile	CGC Arg	ATG Met 275	CAG Gln	GAG Glu	GAG Glu	GAG Glu	AAG Lys 280	AAC Asn	CCC Pro	AAC Asn	ACG Thr	GAG Glu 285	TTC Phe	TAC Tyr	TTG Leu	864
20	AAA Lys	AAC Asn 290	CTG Leu	GTG Val	ATG Met	ACC Thr	ACG Thr 295	TTG Leu	AAC Asn	CTC Leu	TTC Phe	ATT Ile 300	GGG Gly	GGC Gly	ACC Thr	GAG Glu	912
20	ACC Thr 305	GTC Val	AGC Ser	ACC Thr	ACC Thr	CTG Leu 310	CGC Arg	TAT Tyr	GGC Gly	TTC Phe	TTG Leu 315	CTG Leu	CTC Leu	ATG Met	AAG Lys	CAC His 320	960
25	CCA Pro	GAG Glu	GTG Val	GAG Glu	GCC Ala 325	AAG Lys	GTC Val	CAT His	GAG Glu	GAG Glu 330	ATT Ile	GAC Asp	AGA Arg	GTG Val	ATC Ile 335	GGC Gly	1008
20	AAG Lys	AAC Asn	CGG Arg	CAG Gln 340	CCC Pro	AAG Lys	TTT Phe	GAG Glu	GAC Asp 345	CGG Arg	GCC Ala	AAG Lys	ATG Met	CCC Pro 350	TAC Tyr	ATG Met	1056
30															CCC Pro		1104
35															TTC Phe		1152
															CTG Leu		1200
40	GAC Asp	CCC Pro	AGT Ser	TTC Phe	TTC Phe 405	TCC Ser	AAC Asn	CCC Pro	CAG Gln	GAC Asp 410	TTC Phe	AAT Asn	CCC Pro	CAG Gln	CAC His 415	TTC Phe	1248
<b>4</b> 5	CTG Leu	AAT Asn	GAG Glu	AAG Lys 420	GGG Gly	CAG Gln	TTT Phe	AAG Lys	AAG Lys 425	AGT Ser	GAT Asp	GCT Ala	TTT Phe	GTG Val 430	CCC Pro	TTT Phe	1296
	TCC Ser	ATC Ile	GGA Gly 435	AAG Lys	CGG Arg	AAC Asn	TGT Cys	TTC Phe 440	GGA Gly	GAA Glu	GGC Gly	CTG Leu	GCC Ala 445	AGA Arg	ATG Met	GAG Glu	1344

5

			CTC Leu														1392
5	TCC Ser 465	CAG Gln	TCA Ser	CCT Pro	AAG Lys	GAC Asp 470	ATT Ile	GAC Asp	GTG Val	TCC Ser	CCC Pro 475	AGA Arg	CAC His	GTG Val	GGC Gly	TTT Phe 480	1440
10			ATC Ile														1482
	TGA																1485
15	(2)	INFO	OR <b>MA</b> T	rion	FOR	SEQ	ID N	NO: 1	L6:								
		ı	( E	1) LE 3) TY	ENGTH PE:	CHAF H: 49 amir XGY:	94 an	mino cid		_							
20			MOI				_										
		(xi)	SEC	QUENC	CE DE	ESCRI	PTIC	ON: 5	SEQ I	ID NO	): 16	5:					
	Met 1	Leu	Ala	Ser	Gly 5	Met	Leu	Leu	Val	Ala 10	Leu	Leu	Val	Сув	Leu 15	Thr	
25	Val	Met	Val	Leu 20	Met	Ser	Val	Trp	Gln 25	Gln	Arg	Lys	Ser	Lys 30	Gly	Lys	
	Leu	Pro	Pro 35	Gly	Pro	Thr	Pro	Leu 40	Pro	Phe	Ile	Gly	Asn 45	Tyr	Leu	Gln	
30	Leu	Asn 50	Thr	Glu	Gln	Met	Tyr 55	Asn	Ser	Leu	Met	Lys 60	Ile	Ser	Glu	Arg	
	Tyr 65	Gly	Pro	Val	Phe	Thr 70	Ile	His	Leu	Gly	Pro 75	Arg	Arg	Val	Val	Val 80	
35	Leu	Cys	Gly	His	<b>As</b> p 85	Ala	Val	Arg	Glu	Ala 90	Leu	Val	Asp	Gln	Ala 95	Glu	
	Glu	Phe	Ser	Gly 100	Arg	Gly	Glu	Gln	Ala 105	Thr	Phe	Asp	Trp	Val 110	Phe	Lys	
<b>4</b> 0	Gly	Tyr	Gly 115	Val	Val	Phe	Ser	Asn 120	Gly	Glu	Arg	Ala	Lys 125	Gln	Leu	Arg	
	Arg	Phe 130	Ser	Ile	Ala	Thr	Leu 135	Arg	Asp	Phe	Gly	Val 140	Gly	Lys	Arg	Gly	
<b>4</b> 5	Ile 145	Glu	Glu	Arg	Ile	Gln 150	Glu	Glu	Ala	Gly	Phe 155	Leu	Ile	Asp	Ala	Leu 160	
	Arg	Gly	Thr	Gly	Gly 165	Ala	Asn	Ile	Asp	Pro 170	Thr	Phe	Phe	Leu	Ser 175	Arg	

	Thr	Val	Ser	Asn 180	Val	Ile	Ser	Ser	Ile 185	Val	Phe	Gly	Asp	Arg 190	Phe	Asp
5	Tyr	Lys	Asp 195	Lys	Glu	Phe	Leu	Ser 200	Leu	Leu	Arg	Met	Met 205	Leu	Gly	Ile
	Phe	Gln 210	Phe	Thr	Ser	Thr	Ser 215	Thr	Gly	Gln	Leu	Tyr 220	Glu	Met	Phe	Ser
10	Ser 225	Val	Met	Lys	His	Leu 230	Pro	Gly	Pro	Gln	Gln 235	Gln	Ala	Phe	Gln	Leu 240
	Leu	Gln	Gly	Leu	Glu 245	Asp	Phe	Ile	Ala	Lys 250	Lys	Val	Glu	His	Asn 255	Gln
15	Arg	Thr	Leu	Asp 260	Pro	Asn	Ser	Pro	Arg 265	Asp	Phe	Ile	Asp	Ser 270	Phe	Leu
	Ile	Arg	Met 275	Gln	Glu	Glu	Glu	Lys 280	Asn	Pro	Asn	Thr	Glu 285	Phe	Tyr	Leu
20	Lys	Asn 290	Leu	Val	Met	Thr	Thr 295	Leu	Asn	Leu	Phe	Ile 300	Gly	Gly	Thr	Glu
25	Thr 305	Val	Ser	Thr	Thr	Leu 310	Arg	Tyr	Gly	Phe	Leu 315	Leu	Leu	Met	Lys	His 320
	Pro	Glu	Val	Glu	Ala 325	Lys	Val	His	Glu	Glu 330	Ile	Asp	Arg	Val	Ile 335	Gly
30	Lys	Asn	Arg	Gln 340	Pro	Lys	Phe	Glu	Asp 345	Arg	Ala	Lys	Met	Pro 350	Tyr	Met
	Glu	Ala	<b>V</b> al 355	Ile	His	Glu	Ile	Gln 360	Arg	Phe	Gly	Asp	Val 365	Ile	Pro	Met
35	Ser	Leu 370	Ala	Arg	Arg	Val	Lys 375	Lys	Asp	Thr	Lys	Phe 380	Arg	Asp	Phe	Phe
	Leu 385	Pro	Lys	Gly	Thr	Glu 390	Val	Tyr	Pro	Met	Leu 395	Gly	Ser	Val	Leu	Arg 400
<b>4</b> 0	Asp	Pro	Ser	Phe	Phe 405	Ser	Asn	Pro	Gln	Asp 410	Phe	Asn	Pro	Gln	His 415	Phe
	Leu	Asn	Glu	Lys 420	Gly	Gln	Phe	Lys	Lys 425	Ser	Asp	Ala	Phe	Val 430	Pro	Phe
<b>4</b> 5	Ser	Ile	Gly 435	Lys	Arg	Asn	Cys	Phe 440	Gly	Glu	Gly	Leu	Ala 445	Arg	Met	Glu
	Leu	Phe 450	Leu	Phe	Phe	Thr	Thr 455	Val	Met	Gln	Asn	Phe 460	Arg	Leu	Lys	Ser
50	Ser 465	Gln	Ser	Pro	Lys	Asp 470	Ile	Asp	Val	Ser	Pro 475	Arg	His	Val	Gly	Phe 480

Ala	Thr	Ile	Pro	Arg 485	Asn	Tyr	Thr	Met	Ser 490	Phe	Leu	Pro	Arg

(2) INFORMATION FOR SEQ ID NO: 17:

5	(i	(. () ()	A) L: B) T' C) S'	ENGT YPE : TRAN	HARA H: 1- nuc DEDNI OGY:	485 leic ESS:	base aci dou	pai d	rs					
10	(ix	()		AME/I	KEY: ION:		1 <b>4</b> 82							
15	(xi	) SE	QUEN	CE DI	ESCR:	IPTI	ON:	SEQ :	ID N	): 1°	7 :			
					ATG Met									48
20					TCT Ser									96
25					ACC Thr									144
					ATG Met									192
30					ACC Thr 70									240
35					GCC Ala									288
					GGC Gly									336
40					TTC Phe									384
45					ACC Thr									432
					CAG Gln 150									480
50														

53

	CGG Arg	GGC Gly	ACT Thr	GGC Gly	GGC Gly 165	GCC Ala	AAT Asn	ATC Ile	GAT Asp	CCC Pro 170	ACC Thr	TTC Phe	TTC Phe	CTG Leu	AGC Ser 175	CGC Arg	5	528
5		GTC Val										_					5	576
10		AAG Lys															6	24
		CAG Gln 210											_				6	72
15		GTG Val															7	20
20	CTG Leu	CAA Gln	GGG Gly	CTG Leu	GAG Glu 245	GAC Asp	TTC Phe	ATA Ile	GCC Ala	AAG Lys 250	AAG Lys	GTG Val	GAG Glu	CAC His	AAC Asn 255	C <b>A</b> G Gln	7	68
		ACG Thr															8	116
25		CGC Arg															8	64
30		AAC Asn 290												_		_	9	12
		GTC Val															9	60
35		GAG Glu															10	80
<b>4</b> 0		AAC Asn															10	56
40		GCA Ala					Ile						_	_			11	.04
<b>4</b> 5		TTG Leu 370															11	.52
		CCT Pro															12	00

							AAC Asn										1248
5	CTG Leu	AAT Asn	GAG Glu	AAG Lys 420	GGG Gly	CAG Gln	TTT Phe	AAG Lys	AAG Lys 425	AGT Ser	GAT Asp	GCT Ala	TTT Phe	GTG Val 430	CCC Pro	TTT Phe	1296
10							TGT Cys										1344
							ACC Thr 455										1392
15							ATT Ile										1440
20							TAC Tyr										1482
	TGA																1485
25	(2)		(i) S ( <i>I</i> (E	SEQUI A) LE 3) TY	ENCE ENGTI (PE :	CHAF H: 49	ID N RACTE 94 am no ac line	ERIST	rics:	_							
30			MOI	LECUI	E TY	PE:	prot PTIC	ein	SEQ 1	ID NO	): <b>1</b> 8	3:					
	Met 1	Leu	Ala	Ser	Gly 5	Met	Leu	Leu	Val	Ala 10	Leu	Leu	Val	Сув	Leu 15	Thr	
35	Val	Met	Val	Leu 20	Met	Ser	Val	Trp	Gln 25	Gln	Arg	Lys	Ser	Lys 30	Gly	Lys	
	Leu	Pro	Pro 35	Gly	Pro	Thr	Pro	Leu 40	Pro	Phe	Ile	Gly	Asn 45	Tyr	Leu	Gln	
<b>4</b> 0	Leu	Asn 50	Thr	Glu	Gln	Met	Tyr 55	Asn	Ser	Leu	Met	Lys 60	Ile	Ser	Glu	Arg	
	Tyr 65	Gly	Pro	Val	Phe	Thr 70	Ile	His	Leu	Gly	Pro 75	Arg	Arg	Val	Val	Val 80	
<b>4</b> 5	Leu	Суз	Gly	His	Asp 85	Ala	Val	Arg	Glu	Ala 90	Leu	Val	Asp	Gln	Ala 95	Glu	
	Glu	Phe	Ser	Gly 100	Arg	Gly	Glu	Gln	Ala 105	Thr	Phe	Asp	Trp	Val 110	Phe	Lys	
50																	

	Gly	Tyr	Gly 115	Val	Val	Phe	Ser	Asn 120	Gly	Glu	Arg	Ala	Lys 125	Gln	Leu	Arg
5	Arg	Phe 130	Ser	Ile	Ala	Thr	Leu 135	Arg	Asp	Phe	Gly	Val 140	Gly	Lys	Arg	Gly
	Ile 145	Glu	Glu	Arg	Ile	Gln 150	Glu	Glu	Ala	Gly	Phe 155	Leu	Ile	Yab	Ala	Leu 160
10	Arg	Gly	Thr	Gly	Gly 165	Ala	Asn	Ile	Asp	Pro 170	Thr	Phe	Phe	Leu	Ser 175	Arg
	Thr	Val	Ser	Asn 180	Val	Ile	Ser	Ser	Ile 185	Val	Phe	Gly	Asp	Arg 190	Phe	Asp
15	Tyr	Lys	Asp 195	Lys	Glu	Phe	Leu	Ser 200	Leu	Leu	Arg	Met	<b>M</b> et 205	Leu	Gly	Ile
20	Phe	Gln 210	Phe	Thr	Ser	Thr	Ser 215	Thr	Gly	Gln	Leu	Tyr 220	Glu	Met	Phe	Ser
20	Ser 225	Val	Met	Lys	His	Leu 230	Pro	Gly	Pro	Gln	Gln 235	Gln	Ala	Phe	Gln	Leu 240
25	Leu	Gln	Gly	Leu	Glu 245	Asp	Phe	Ile	Ala	Lys 250	Lys	Val	Glu	His	<b>A</b> sn 255	Gln
	Arg	Thr	Leu	Asp 260	Pro	Asn	Ser	Pro	Arg 265	Asp	Phe	Ile	Asp	Ser 270	Phe	Leu
30	Ile	Arg	Met 275	Gln	Glu	Glu	Glu	Lys 280	Asn	Pro	Asn	Thr	Glu 285	Phe	Tyr	Leu
	Lys	Asn 290	Leu	Val	Met	Thr	Thr 295	Leu	Asn	Leu	Phe	Ile 300	Gly	Gly	Thr	Glu
35	Thr 305	Val	Ser	Thr	Thr	Leu 310	Arg	Tyr	Gly	Phe	Leu 315	Leu	Leu	Met	Lys	His 320
	Pro	Glu	Val	Glu	Ala 325	Lys	Val	His	Glu	Glu 330	Ile	Asp	Arg	Val	Ile 335	Gly
<b>4</b> 0	Lys	Asn	Arg	Gln 3 <b>4</b> 0	Pro	Lys	Phe	Glu	Asp 345	Arg	Ala	Lys	Met	Pro 350	Tyr	Met
	Glu	Ala	<b>Val</b> 355	Ile	His	Glu	Ile	Gln 360	Arg	Phe	Gly	Asp	<b>Val</b> 365	Ile	Pro	Met
<b>4</b> 5	Ser	Leu 370	Ala	Arg	Arg	Val	Lys 375	Lys	Asp	Thr	Lys	Phe 380	Arg	Asp	Phe	Phe
	Leu 385	Pro	Lys	Gly	Thr	Glu 390	Val	Tyr	Pro	Met	Leu 395	Gly	Ser	Val	Leu	Arg 400
50	Asp	Pro	Ser	Phe	Phe 405	Ser	Asn	Pro	Gln	Asp 410	Phe	Asn	Pro	Gln	His 415	Phe

	Leu	Asn	Glu	Lys 420	Gly	Gln	Phe	Lys	Lys 425	Ser	Asp	Ala	Phe	Val 430	Pro	Phe	
5	Ser	Ile	Gly 435	Lys	Arg	Asn	Cys	Phe 440	Gly	Glu	Gly	Leu	Ala 445	Arg	Met	Glu	
	Leu	Phe 450	Leu	Phe	Phe	Thr	Thr 455	Val	Met	Gln	Asn	Phe 460	Arg	Leu	Lys	Ser	
10	Ser 465	Gln	Ser	Pro	Lys	Asp 470	Ile	Asp	Val	Ser	Pro 475	Lys	His	Val	Gly	Phe 480	
	Ala	Thr	Ile	Pro	Arg 485	Asn	Tyr	Thr	Met	Ser 490	Phe	Leu	Pro	Arg			
	(2)	INF	ORMAT	rion	FOR	SEQ	ID 1	<b>1</b> O: 1	19:								
15		(i)	() (E	QUENCA) LE B) TY C) ST C) TO	ENGTI (PE : [R <b>AN</b> I	H: 14 nucl	176 h Leic ESS:	ase acio doub	pain i	cs							
20		(ix)	(1	ATURI A) <b>N</b> A 3) LO	ME/I			473									
25		(xi)	SEÇ	QUENC	CE DE	ESCRI	PTIC	ON: 5	SEQ 1	D NO	): 19	<b>)</b> :					
				AGC Ser													4
30	CTA Leu	CTC Leu	CTG Leu	GTT Val 20	CAG Gln	CGC <b>A</b> rg	CAC His	CCT Pro	AAC Asn 25	ACC Thr	CAT His	GAC Asp	CGC Arg	CTC Leu 30	CCA Pro	CCA Pro	9
35	GGG Gly	CCC Pro	CGC Arg 35	CCT Pro	CTG Leu	CCC Pro	CTT Leu	TTG Leu 40	GGA Gly	AAC Asn	CTT Leu	CTG Leu	CAG Gln 45	ATG Met	GAT Asp	AGA Arg	14
	AGA Arg	GGC Gly 50	CTA Leu	CTC Leu	AAA Lys	TCC Ser	TTT Phe 55	CTG Leu	AGG Arg	TTC Phe	CGA Arg	GAG Glu 60	AAA Lys	тат Туг	GGG Gly	GAC Asp	19
<b>4</b> 0	GTC Val 65	TTC Phe	ACG Thr	GTA Val	CAC His	CTG Leu 70	GGA Gly	CCG Pro	AGG Arg	CCC Pro	GTG Val 75	GTC Val	ATG Met	CTG Leu	TGT Cys	GGA Gly 80	24
<b>4</b> 5	GTA Val	GAG Glu	GCC Ala	ATA Ile	CGG Arg 85	GAG Glu	GCC Ala	CTT Leu	GTG Val	GAC Asp 90	AAG Lys	GCT Ala	GAG Glu	GCC Ala	TTC Phe 95	TCT Ser	28
	GGC Gly	CGG <b>A</b> rg	GGA Gly	AAA Lys 100	ATC Ile	GCC Ala	ATG Met	GTC Val	GAC Asp 105	CCA Pro	TTC Phe	TTC Phe	CGG Arg	GGA Gly 110	TAT Tyr	GGT Gly	33

	GTG Val	ATC Ile	TTT Phe 115	GCC Ala	AAT Asn	GGA Gly	AAC Asn	CGC Arg 120	TGG Trp	AAG Lys	GTG Val	CTT Leu	CGG Arg 125	CGA Arg	TTC Phe	TCT Ser	3	38 <b>4</b>
5	GTG Val	ACC Thr 130	ACT Thr	ATG Met	AGG Arg	GAC Asp	TTC Phe 135	GGG Gly	ATG Met	GGA Gly	AAG Lys	CGG Arg 140	AGT Ser	GTG Val	GAG Glu	GAG Glu	4	132
10	CGG Arg 145	ATT Ile	CAG Gln	GAG Glu	GAG Glu	GCT Ala 150	CAG Gln	TGT Cys	CTG Leu	ATA Ile	GAG Glu 155	GAG Glu	CTT Leu	CGG Arg	AAA Lys	TCC Ser 160	4	180
	AAG Lys	GGG Gly	GCC Ala	CTC Leu	ATG Met 165	GAC Asp	CCC Pro	ACC Thr	TTC Phe	CTC Leu 170	TTC Phe	CAG Gln	TCC Ser	ATT Ile	ACC Thr 175	GCC Ala	Ē	528
15	AAC Asn	ATC Ile	ATC Ile	TGC Cys 180	TCC Ser	ATC Ile	GTC Val	TTT Phe	GGA Gly 185	AAA Lys	CGA Arg	TTC Phe	CAC His	TAC Tyr 190	CAA Gln	GAT Asp	<u>5</u>	576
20	CAA Gln	GAG Glu	TTC Phe 195	CTG Leu	AAG Lys	ATG Met	CTG Leu	AAC Asn 200	TTG Leu	TTC Phe	TAC Tyr	CAG Gln	ACT Thr 205	TTT Phe	TCA Ser	CTC Leu	6	524
	ATC Ile	AGC Ser 210	TCT Ser	GTA Val	TTC Phe	GGC Gly	CAG Gln 215	CTG Leu	TTT Phe	GAG Glu	CTC Leu	TTC Phe 220	TCT Ser	GGC Gly	TTC Phe	TTG Leu	6	572
25	AAA Lys 225	TAC Tyr	TTT Phe	CCT Pro	GGG Gly	GCA Ala 230	CAC His	AGG Arg	CAA Gln	GTT Val	TAC Tyr 235	AAA Lys	AAC Asn	CTG Leu	CAG Gln	GAA Glu 240	7	720
30	ATC Ile	AAT Asn	GCT Ala	TAC Tyr	ATT Ile 245	GGC Gly	CAC His	AGT Ser	GTG Val	GAG Glu 250	AAG Lys	CAC His	CGT Arg	GAA Glu	ACC Thr 255	CTG Leu	7	768
	GAC Asp	CCC Pro	AGC Ser	GCC Ala 260	CCC Pro	AAG Lys	GAC Asp	CTC Leu	ATC Ile 265	GAC Asp	ACC Thr	TAC Tyr	CTG Leu	CTC Leu 270	CAC His	ATG Met	ε	316
35	GAA Glu	AAA Lys	GAG Glu 275	AAA Lys	TCC Ser	AAC Asn	GCA Ala	CAC His 280	AGT Ser	GAA Glu	TTC Phe	AGC Ser	CAC His 285	CAG Gln	AAC Asn	CTC Leu	ε	364
<b>4</b> 0	AAC Asn	CTC Leu 290	AAC Asn	ACG Thr	CTC Leu	TCG Ser	CTC Leu 295	TTC Phe	TTT Phe	GCT Ala	GGC Gly	ACT Thr 300	GAG Glu	ACC Thr	ACC Thr	AGC Ser	g	912
	ACC Thr 305	ACT Thr	CTC Leu	CGC <b>A</b> rg	TAC Tyr	GGC Gly 310	TTC Phe	CTG Leu	CTC Leu	Met	CTC Leu 315	Lys	TAC Tyr	CCT Pro	CAT His	GTT Val 320	Ş	960
<b>4</b> 5	GCA Ala	GAG Glu	AGA Arg	GTC Val	TAC Tyr 325	AGG Arg	GAG Glu	ATT Ile	GAA Glu	CAG Gln 330	GTG Val	ATT Ile	GGC Gly	CCA Pro	CAT His 335	CGC <b>A</b> rg	10	800
50	CCT Pro	CCA Pro	GAG Glu	CTT Leu 340	CAT His	GAC <b>A</b> sp	CGA <b>A</b> rg	GCC <b>A</b> la	AAA Lys 345	ATG Met	CCA Pro	TAC Tyr	ACA Thr	GAG Glu 350	GCA Ala	GTC Val	10	056

		TAT Tyr															1104
5	CAC His	ATT Ile 370	GTC Val	ACC Thr	C <b>AA</b> Gln	CAC His	ACC Thr 375	AGC Ser	TTC Phe	CGA Arg	GGG Gly	TAC Tyr 380	ATC Ile	ATC Ile	CCC Pro	AAG Lys	1152
10	GAC Asp 385	ACA Thr	GAA Glu	GTA Val	TTT Phe	CTC Leu 390	ATC Ile	CTG Leu	AGC Ser	ACT Thr	GCT Ala 395	CTC Leu	CAT His	GAC Asp	CCA Pro	CAC His 400	1200
		TTT Phe															1248
15		GGG Gly															1296
20	AAG Lys	CGG Arg	ATT Ile 435	TGT Cys	CTT Leu	GGT Gly	GAA Glu	GGC Gly 440	ATC Ile	GCC <b>A</b> la	CGT <b>A</b> rg	GCG Ala	GAA Glu 445	TTG Leu	TTC Phe	CTC Leu	1344
		TTC Phe 450															1392
25	CCA Pro 465	G <b>AA</b> Glu	GAC Asp	ATC Ile	GAT Asp	CTG Leu 470	ACA Thr	CCC Pro	CAG Gln	GAG Glu	TGT Cys 475	GGT Gly	GTG Val	GGC Gly	AAA Lys	ATA Ile 480	1440
30		CCA Pro										TGA					1476
	(2)	INFO	RMAT	ION	FOR	SEQ	ID N	IO: 2	0:								
35		(	( E	L) LE	ENCE ENGTH PE: POLC	I: 49 amir	01 am	ino id									
		(ii)	MOL	ECUI	E TY	PE:	prot	ein									
<b>\$</b> 0		(xi)	SEÇ	UENC	CE DE	SCRI	PTIC	N: S	EQ I	D <b>N</b> C	): 20	);					
-	Met 1	Glu	Leu	Ser	Val 5	Leu	Leu	Phe	Leu	Ala 10	Leu	Leu	Thr	Gly	Leu 15	Leu	
	Leu	Leu	Leu	Val 20	Gln	Arg	His	Pro	Asn 25	Thr	His	Asp	Arg	Leu 30	Pro	Pro	

59

Gly Pro Arg Pro Leu Pro Leu Leu Gly Asn Leu Leu Gln Met Asp Arg 35 40 45

50

	Arg	Gly 50	Leu	Leu	Lys	Ser	Phe 55	Leu	Arg	Phe	Arg	Glu 60	Lys	Tyr	Gly	Asp
5	Val 65	Phe	Thr	Val	His	Leu 70	Gly	Pro	Arg	Pro	Val 75	Val	Met	Leu	Cys	Gly 80
	Val	Glu	Ala	Ile	Arg 85	Glu	Ala	Leu	Val	<b>A</b> sp 90	Lys	Ala	Glu	Ala	Phe 95	Ser
10	Gly	Arg	Gly	Lys 100	Ile	Ala	Met	Val	<b>Asp</b> 105	Pro	Phe	Phe	Arg	Gly 110	Tyr	Gly
	Val	Ile	Phe 115	Ala	Asn	Gly	Asn	Arg 120	Trp	Lys	Val	Leu	Arg 125	Arg	Phe	Ser
15	Val	Thr 130	Thr	Met	Arg	Asp	Phe 135	Gly	Met	Gly	Lys	Arg 140	Ser	Val	Glu	Glu
	Arg 145	Ile	Gln	Glu	Glu	Ala 150	Gln	Cys	Leu	Ile	Glu 155	Glu	Leu	Arg	Lys	Ser 160
20	Lys	Gly	Ala	Leu	Met 165	Asp	Pro	Thr	Phe	Leu 170	Phe	Gln	Ser	Ile	Thr 175	Ala
	Asn	Ile	Ile	Cys 180	Ser	Ile	Val	Phe	Gly 185	Lys	Arg	Phe	His	Tyr 190	Gln	Asp
25	Gln	Glu	Phe 195	Leu	Lys	Met	Leu	Asn 200	Leu	Phe	Tyr	Gln	Thr 205	Phe	Ser	Leu
		210					215					220		Gly		
30	Lys 225	Tyr	Phe	Pro	Gly	Ala 230	His	Arg	Gln	Val	Tyr 235	Lys	Asn	Leu	Gln	Glu 2 <b>4</b> 0
					<b>24</b> 5					250				Glu	255	
35	Asp	Pro	Ser	Ala 260	Pro	Lys	Asp	Leu	Ile 265	Asp	Thr	Tyr	Leu	Leu 270	His	Met ,
			<b>27</b> 5					280					285	Gln		
40	Asn	Leu 290	Asn	Thr	Leu	Ser	Leu 295	Phe	Phe	Ala	Gly	Thr 300	Glu	Thr	Thr	Ser
	Thr 305	Thr	Leu	Arg	Tyr	Gly 310	Phe	Leu	Leu	Met	Leu 315	Lys	Tyr	Pro	His	Val 320
45	Ala	Glu	Arg	Val	Tyr 325	Arg	Glu	Ile	Glu	Gln 330	Val	Ile	Gly	Pro	His 335	Arg
	Pro	Pro	Glu	Leu 340	His	Asp	Arg	Ala	Lys 345	Met	Pro	Tyr	Thr	Glu 350	Ala	Val
50	Ile	Tyr	Glu 355	Ile	Gln	Arg	Phe	Ser 360	Asp	Leu	Leu	Pro	<b>Me</b> t 365	Gly	Val	Pro

	His	Ile 370	Val	Thr	Gln	His	Thr 375	Ser	Phe	Arg	Gly	Tyr 380	Ile	Ile	Pro	Lys	
5	Asp 385	Thr	Glu	Val	Phe	Leu 390	Ile	Leu	Ser	Thr	Ala 395	Leu	His	Asp	Pro	His 400	
	Tyr	Phe	Glu	Lys	Pro 405	Asp	Ala	Phe	Asn	Pro 410	Asp	His	Phe	Leu	Asp 415	Ala	
10	Asn	Gly	Ala	Leu 420	Lys	Lys	Thr	Glu	Ala 425	Phe	Ile	Pro	Phe	Ser 430	Leu	Gly	
	Lys	Arg	Ile 435	Cys	Leu	Gly	Glu	Gly 440	Ile	Ala	Arg	Ala	Glu 445	Leu	Phe	Leu	
15	Phe	Phe 450	Thr	Thr	Ile	Leu	Gln <b>4</b> 55	Asn	Phe	Ser	Met	Ala 460	Ser	Pro	Val	Ala	
	Pro 465	Glu	Asp	Ile	Asp	Leu 470	Thr	Pro	Gln	Glu	Cys 475	Gly	Val	Gly	Lys	Ile 480	
20	Pro	Pro	Thr	Tyr	Gln 485	Ile	Arg	Phe	Leu	Pro 490	Arg						
	(2)		ORMAT SEÇ														
25		(1)	( F ( E	A) LE B) TY C) ST	ENGTH PE: TRANI	I: 14 nucl	73 h leic ESS: line	ase acio doub	pair l_	្ន							
30		(ix)		A) NA	ME/F	ŒY: ON:	CDS 11	470									
			SEC										tanta.	N TO C	Omm	CTC	4.0
35	ATG Met	GAA	SEÇ CCT Pro	ттт	GTG	GTC	CTG	GTG	CTG	TGT	CTC	TCT	TTT Phe	ATG Met	CTT Leu 15	CTC Leu	48
35	Met 1 TTT	GAA Glu TCA	CCT	TTT Phe	GTG Val 5	GTC Val	CTG Leu AGC	GTG Val	CTG Leu AGG	TGT Cys 10	CTC Leu AGG	TCT Ser	Phe CTC	Met CCT	Leu 15 CCT	Leu GGC	<b>48</b> 96
35 40	Met 1 TTT Phe	GAA Glu TCA Ser	CCT Pro	TTT Phe TGG Trp 20 CTT	GTG Val 5 AGA Arg	GTC Val CAG Gln	CTG Leu AGC Ser	GTG Val TGT Cys	CTG Leu AGG Arg 25	TGT Cys 10 AGA Arg	CTC Leu AGG Arg	TCT Ser AAG Lys	Phe CTC Leu ATA	Met CCT Pro 30 GAT	Leu 15 CCT Pro	GGC Gly	
	Met 1 TTT Phe CCC Pro	GAA Glu TCA Ser ACT Thr	CCT Pro	TTT Phe TGG Trp 20 CTT Leu	GTG Val 5 AGA Arg CCT Pro	GTC Val CAG Gln ATT Ile	CTG Leu AGC Ser ATT Ile	GTG Val TGT Cys GGA Gly 40	CTG Leu AGG Arg 25 AAT Asn	TGT Cys 10 AGA Arg ATG Met	CTC Leu AGG Arg CTA Leu	TCT Ser AAG Lys CAG Gln	Phe CTC Leu ATA Ile 45	Met CCT Pro 30 GAT Asp	Leu 15 CCT Pro GTT Val	GGC Gly AAG Lys	96
<b>4</b> 0	Met 1 TTT Phe CCC Pro	GAA Glu TCA Ser ACT Thr	CCT Pro CTC Leu CCT Pro 35	TTT Phe TGG Trp 20 CTT Leu	GTG Val 5 AGA Arg CCT Pro	GTC Val CAG Gln ATT Ile	CTG Leu AGC Ser ATT Ile ACC Thr	GTG Val TGT Cys GGA Gly 40	CTG Leu AGG Arg 25 AAT Asn	TGT Cys 10 AGA Arg ATG Met	CTC Leu AGG Arg CTA Leu	TCT Ser AAG Lys CAG Gln GTC Val	Phe CTC Leu ATA Ile 45	Met CCT Pro 30 GAT Asp	Leu 15 CCT Pro GTT Val	GGC Gly AAG Lys	96 144

	TTC Phe 65	ACC Thr	GTG Val	TAT Tyr	TTT Phe	GGC Gly 70	ATG Met	AAT Asn	CCC Pro	ATA Ile	GTG Val 75	GTG Val	TTT Phe	CAT His	GGA Gly	TAT Tyr 80	240
5	GAG Glu	GCA Ala	GTG Val	AAG Lys	GAA Glu 85	GCC Ala	CTG Leu	ATT Ile	GAT Asp	AAT Asn 90	GGA Gly	GAG Glu	GAG Glu	TTT Phe	TCT Ser 95	GGA Gly	288
10	AGA Arg	GGC Gly	AAT Asn	TCC Ser 100	CCA Pro	ATA Ile	TCT Ser	CAA Gln	AGA Arg 105	ATT Ile	ACT Thr	AAA Lys	GGA Gly	CTT Leu 110	GGA Gly	ATC Ile	336
	ATT Ile	TCC Ser	AGC Ser 115	AAT Asn	GGA Gly	AAG Lys	AGA Arg	TGG Trp 120	AAG Lys	GAG Glu	ATC Ile	CGG Arg	CGT Arg 125	TTC Phe	TCC Ser	CTC Leu	384
15	ACA Thr	ACC Thr 130	TTG Leu	CGG <b>A</b> rg	AAT Asn	TTT Phe	GGG Gly 135	ATG Met	GGG Gly	AAG Lys	AGG Arg	AGC Ser 140	ATT Ile	GAG Glu	GAC Asp	CGT Arg	432
20	GTT Val 145	C <b>AA</b> Gln	GAG Glu	GAA Glu	GCT Ala	CAC His 150	TGC Cys	CTT Leu	GTG Val	GAG Glu	GAG Glu 155	TTG Leu	AGA Arg	AAA Lys	ACC Thr	AAG Lys 160	480
	GCT Ala	TCA Ser	CCC Pro	TGT Cys	GAT Asp 165	CCC Pro	ACT Thr	TTC Phe	ATC Ile	CTG Leu 170	GGC Gly	TGT Cys	GCT Ala	CCC Pro	TGC Cys 175	AAT Asn	528
25	GTG Val	ATC Ile	TGC Cys	TCC Ser 180	GTT Val	GTT Val	TTC Phe	CAG Gln	AAA Lys 185	CGA Arg	TTT Phe	GAT Asp	TAT Tyr	AAA Lys 190	GAT Asp	CAG Gln	576
30	Asn	Phe	Leu 195	Thr	Leu	Met	Lys	Arg 200	TTC Phe	Asn	Glu	Asn	Phe 205	Arg	Ile	Leu	624
	Asn	Ser 210	Pro	Trp	Ile	Gln	Val 215	Cys	AAT Asn	Asn	Phe	Pro 220	Leu	Leu	Ile	Asp	672
35	TGT Cys 225	TTC Phe	CCA Pro	GGA Gly	ACT Thr	CAC His 230	AAC Asn	AAA Lys	GTG Val	CTT Leu	AAA Lys 235	AAT Asn	GTT Val	GCT Ala	CTT Leu	ACA Thr 240	720
40	Arg	Ser	Tyr	Ile	<b>Ar</b> g 2 <b>4</b> 5	Glu	Lys	Val	AAA Lys	Glu 250	His	Gln	Ala	Ser	Leu 255	Asp	768
	GTT Val	AAC Asn	Asn	CCT Pro 260	Arg	GAC Asp	Phe	Ile	GAT Asp 265	Cys	TTC Phe	Leu	Ile	Lys	Met	GAG Glu	816
<b>4</b> 5									G <b>AA</b> Glu								864
50	GGC Gly	ACT Thr 290	Val	GCT <b>A</b> la	GAT Asp	CTA Leu	TTT Phe 295	GTT Val	GCT Ala	GGA Gly	ACA Thr	GAG Glu 300	ACA Thr	ACA Thr	AGC Ser	ACC Thr	912

		-	. ~ .			om a	ama	ama	CTIC	CTC	220	CNC	CCA	CAC	CTC	A C'A	960
	ACT Thr 305	CTG Leu	AGA Arg	TAT	GGA	CTC Leu 310	Leu	Leu	Leu	Leu	Lys 315	His	Pro	Glu	Val	Thr 320	960
5	GCT Ala	AAA Lys	GTC Val	CAG Gln	GAA Glu 325	GAG Glu	ATT Ile	GAT Asp	CAT His	GTA Val 330	ATT Ile	GGC Gly	AGA Arg	CAC His	AGG Arg 335	AGC Ser	1008
10	CCC Pro	TGC Cys	ATG Met	CAG Gln 340	GAT Asp	AGG Arg	AGC Ser	CAC His	ATG Met 345	CCT Pro	TAC Tyr	ACT Thr	GAT Asp	GCT Ala 350	GTA Val	GTG Val	1056
	CAC His	GAG Glu	ATC Ile 355	CAG Gln	AGA Arg	TAC Tyr	AGT Ser	GAC Asp 360	CTT Leu	GTC Val	CCC Pro	ACC Thr	GGT Gly 365	GTG Val	CCC Pro	CAT His	1104
15	GCA Ala	GTG Val 370	ACC Thr	ACT Thr	GAT Asp	ACT Thr	AAG Lys 375	TTC Phe	AGA Arg	AAC Asn	TAC Tyr	CTC Leu 380	ATC Ile	CCC Pro	AAG Lys	GGC Gly	1152
20	ACA Thr 385	ACC Thr	ATA Ile	ATG Met	GCA Ala	TTA Leu 390	CTG Leu	ACT Thr	TCC Ser	GTG Val	CTA Leu 395	CAT His	GAT Asp	GAC Asp	AAA Lys	GAA Glu 400	1200
	TTT Phe	CCT Pro	AAT Asn	CCA Pro	AAT Asn 405	ATC Ile	TTT Phe	GAC Asp	CCT Pro	GGC Gly 410	CAC His	TTT Phe	CTA Leu	GAT Asp	AAG Lys 415	AAT Asn	1248
25	GGC Gly	AAC Asn	TTT Phe	AAG Lys 420	AAA Lys	AGT Ser	GAC Asp	TAC Tyr	TTC Phe 425	ATG Met	CCT Pro	TTC Phe	TCA Ser	GCA Ala 430	GGA Gly	AAA Lys	1296
30	CGA Arg	ATT Ile	TGT Cys 435	GCA Ala	GGA Gly	GAA Glu	GGA Gly	CTT Leu 440	GCC Ala	CGC Arg	ATG Met	GAG Glu	CTA Leu 445	TTT Phe	TTA Leu	TTT Phe	1344
	CTA Leu	ACC Thr 450	ACA Thr	ATT Ile	TTA Leu	CAG Gln	AAC Asn 455	TTT Phe	AAC Asn	CTG Leu	AAA Lys	TCT Ser 460	GTT Val	GAT Asp	GAT Asp	TTA Leu	1392
35	AAG Lys 465	AAC Asn	CTC Leu	AAT Asn	ACT Thr	ACT Thr 470	GCA Ala	GTT Val	ACC Thr	AAA Lys	GGG Gly 475	ATT Ile	GTT Val	TCT Ser	CTG Leu	CCA Pro 480	1440
<b>1</b> 0						TGC Cys					TGA						1473

(2) INFORMATION FOR SEQ ID NO: 22:

(i) SEQUENCE CHARACTERISTICS:
(A) LENGTH: 490 amino acids
(B) TYPE: amino acid
(D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

55

45

# (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 22:

5	Met 1	Glu	Pro	Phe	Val 5	Val	Leu	Val	Leu	Cys 10	Leu	Ser	Phe	Met	Leu 15	Leu
	Phe	Ser	Leu	Trp 20	Arg	Gln	Ser	Cys	Arg 25	Arg	Arg	Lys	Leu	Pro 30	Pro	Gly
10	Pro	Thr	Pro 35	Leu	Pro	Ile	Ile	Gly 40	Asn	Met	Leu	Gln	Ile 45	Asp	Val	Lys
	Asp	Ile 50	Cys	Lys	Ser	Phe	Thr 55	Asn	Phe	Ser	Lys	Val 60	Tyr	Gly	Pro	Val
15	Phe 65	Thr	Val	Tyr	Phe	Gly 70	Met	Asn	Pro	Ile	Val 75	Val	Phe	His	Gly	Tyr 80
	Glu	Ala	Val	Lys	Glu 85	Ala	Leu	Ile	Asp	Asn 90	Gly	Glu	Glu	Phe	Ser 95	Gly
20	Arg	Gly	Asn	Ser 100	Pro	Ile	Ser	Gln	Arg 105	Ile	Thr	Lys	Gly	Leu 110	Gly	Ile
	Ile	Ser	Ser 115	Asn	Gly	Lys	Arg	Trp 120	Lys	Glu	Ile	Arg	Arg 125	Phe	Ser	Leu
25		Thr 130	Leu	Arg	Asn	Phe	Gly 135	Met	Gly	Lys	Arg	Ser 140	Ile	Glu	Asp	Arg
	Val 145	Gln	Glu	Glu	Ala	His 150	Cys	Leu	Val	Glu	Glu 155	Leu	Arg	Lys	Thr	Lys 160
30	Ala	Ser	Pro	Cys	Asp 165	Pro	Thr	Phe	Ile	Leu 170	Gly	Суз	Ala	Pro	Суs 175	Asn
	Val	Ile	Cys	Ser 180	Val	Val	Phe	Gln	Lys 185	Arg	Phe	Asp	Tyr	Lys 190	Asp	Gln
35	Asn	Phe	Leu 195	Thr	Leu	Met	Lys	Arg 200	Phe	Asn	Glu	Asn	Phe 205	Arg	Ile	Leu
	Asn	Ser 210	Pro	Trp	Ile	Gln	Val 215	Cys	Asn	Asn	Phe	Pro 220	Leu	Leu	Ile	Ąsp
<b>4</b> 0	Cys 225	Phe	Pro	Gly	Thr	His 230	Asn	Lys	Val	Leu	Lys 235	Asn	Val	Ala	Leu	Thr 240
	Arg	Ser	Tyr	Ile	Arg 245	Glu	Lys	Val	Lys	Glu 250	His	Gln	Ala	Ser	Leu 255	Asp
45	Val	Asn	Asn	Pro 260	Arg	Asp	Phe	Ile	<b>Asp</b> 265	Cys	Phe	Leu	Ile	Lys 270	Met	Glu
	Gln	Glu	Lys 275	Asp	Asn	Gln	Lys	Ser 280	Glu	Phe	Asn	Ile	Glu 285	Asn	Leu	Val
50	Gly	Thr 290	Val	Ala	Asp	Leu	Phe 295	Val	Ala	Gly	Thr	Glu 300	Thr	Thr	Ser	Thr

	Thr 305	Leu	Arg	Tyr	Gly	Leu 310	Leu	Leu	Leu	Leu	Lys 315	His	Pro	Glu	Val	Thr 320	
5	Ala	Lys	Val	Gln	Glu 325	Glu	Ile	Asp	His	Val 330	Ile	Gly	Arg	His	<b>A</b> rg 335	Ser	
	Pro	Cys	Met	Gln 340	Asp	Arg	Ser	His	Met 345	Pro	Tyr	Thr	Asp	Ala 350	Val	Val	
10	His	Glu	Ile 355	Gln	Arg	Tyr	Ser	Asp 360	Leu	Val	Pro	Thr	Gly 365	Val	Pro	His	
	Ala	Val 370	Thr	Thr	Asp	Thr	Lys 375	Phe	Arg	Asn	Tyr	Leu 380	Ile	Pro	Lys	Gly	
15	Thr 385	Thr	Ile	Met	Ala	Leu 390	Leu	Thr	Ser	Val	Leu 395	His	Asp	Asp	Lys	Glu 400	
	Phe	Pro	Asn	Pro	Asn 405	Ile	Phe	Asp	Pro	Gly 410	His	Phe	Leu	Asp	Lys 415	Asn	
20	Gly	Asn	Phe	Lys 420	Lys	Ser	Asp	Tyr	Phe 425	Met	Pro	Phe	Ser	Ala 430	Gly	Lys	
	Arg	Ile	Cys 435	Ala	Gly	Glu	Gly	Leu 440	Ala	Arg	Met	Glu	Leu 445	Phe	Leu	Phe	
25	Leu	Thr 450	Thr	Ile	Leu	Gln	<b>As</b> n <b>4</b> 55	Phe	Asn	Leu	Lys	Ser 460	Val	Asp	Asp	Leu	
	Lys 465	Asn	Leu	Asn	Thr	Thr 470	Ala	Val	Thr	Lys	Gly 475	Ile	Val	Ser	Leu	Pro 480	
30	Pro	Ser	Tyr	Gln	Ile 485	Суѕ	Phe	Ile	Pro	Val 490							
	(2)	INFO	RMAT	CION	FOR	SEQ	ID N	IO: 2	?3 :								
35		(i)	(E	UENCAL LE LE LE LE CE LE LE CE	ENGTE PE: TRANI	I: 14 nucl EDNE	73 b eic ESS:	ase acid doub	pair 1	rs							
<b>4</b> 0		(ix)		TURE A) NA B) LC	ME/K			470									
		(xi)	SEÇ	UENC	E DE	SCRI	PTIC	N: 5	SEQ I	D NC	): 23	3:					
<b>4</b> 5			CCT Pro														4.8
5 <i>0</i>			CTC Leu														91

			ATT						144
5			TTC Phe						192
10			GGC Gly 70						240
			GCC Ala						288
15			ATA Ile						336
20			AAG Lys						384
20			TTT Phe						432
25			CAC His 150						480
20			CCC Pro						528
30			GTT Val						576
35			ATG Met						624
			CAG Gln						672
<b>4</b> 0			CAC His 230						720
<b>4</b> 5			GAG Glu						768

	GTT Val	AAC Asn	AAT Asn	CCT Pro 260	CGG Arg	GAC Asp	TTT Phe	ATC Ile	GAT Asp 265	ТGC	TTC Phe	CTG Leu	ATC Ile	AAA Lys 270	ATG Met	GAG Glu	816
5	CAG Gln	G <b>AA</b> Glu	AAG Lys 275	GAC Asp	AAC Asn	C <b>AA</b> Gln	AAG Lys	TCA Ser 280	GAA Glu	TTC Phe	AAT Asn	ATT Ile	GAA Glu 285	AAC Asn	TTG Leu	GTT Val	864
10	GGC Gly	ACT Thr 290	GTA Val	GCT Ala	GAT <b>A</b> sp	CTA Leu	TTT Phe 295	GTT Val	GCT Ala	GGA Gly	ACA Thr	GAG Glu 300	ACA Thr	ACA Thr	AGC Ser	ACC Thr	912
	ACT Thr 305	CTG Leu	AGA Arg	TAT Tyr	GGA Gly	CTC Leu 310	CTG Leu	CTC Leu	CTG Leu	CTG Leu	AAG Lys 315	CAC His	CCA Pro	GAG Glu	GTC Val	ACA Thr 320	960
15	GCT Ala	AAA Lys	GTC Val	CAG Gln	GAA Glu 325	G <b>A</b> G Glu	ATT Ile	GAT Asp	CAT His	GTA Val 330	ATT Ile	GGC Gly	AGA Arg	CAC His	AGG Arg 335	AGC Ser	1008
20	CCC Pro	TGC Cys	ATG Met	CAG Gln 340	GAT Asp	AGG Arg	AGC Ser	CAC His	ATG Met 345	CCT Pro	TAC Tyr	ACT Thr	GAT Asp	GCT Ala 350	GTA Val	GTG Val	1056
	CAC His	GAG Glu	ATC Ile 355	CAG Gln	AGA Arg	TAC Tyr	AGT Ser	GAC Asp 360	CTT Leu	GTC Val	CCC Pro	ACC Thr	GGT Gly 365	GTG Val	CCC Pro	CAT His	1104
25	GCA Ala	GTG Val 370	ACC Thr	ACT Thr	GAT Asp	ACT Thr	AAG Lys 375	TTC Phe	AGA Arg	AAC Asn	TAC Tyr	CTC Leu 380	ATC Ile	CCC Pro	AAG Lys	GGC Gly	1152
20						TTA Leu 390											1200
30	TTT Phe	CCT Pro	AAT Asn	CCA Pro	AAT Asn 405	ATC Ile	TTT Phe	GAC Asp	CCT Pro	GGC Gly 410	CAC His	TTT Phe	CTA Leu	GAT Asp	AAG Lys 415	AAT Asn	1248
35	GGC Gly	AAC Asn	TTT Phe	AAG Lys 420	AAA Lys	AGT Ser	GAC Asp	TAC Tyr	TTC Phe 425	ATG Met	CCT Pro	TTC Phe	TCA Ser	GCA Ala 430	GGA Gly	AAA Lys	1296
						GAA Glu											1344
40	CTA Leu	ACC Thr 450	ACA Thr	ATT Ile	TTA Leu	CAG Gln	AAC Asn 455	TTT Phe	AAC Asn	CTG Leu	AAA Lys	TCT Ser 460	GTT Val	GAT Asp	GAT Asp	TTA Leu	1392
<b>4</b> 5						ACT Thr 470											1440
						TGC Cys					TGA						1473

	(2)	INF	OR <b>MA</b> T	rion	FOR	SEQ	ID i	10: 2	24:							
5			(I	A) LI 3) TY	ENGTI PE:	H: 49 amir		mino cid								
		(ii)	MOI	LECUI	LE TY	PE:	prot	cein								
10		(xi)	) SE(	QUEN	CE DE	ESCRI	IPTIC	ON: S	SEQ I	D N	): 24	1:				
	Met 1	Glu	Pro	Phe	Val 5	Val	Leu	Val	Leu	Cys 10	Leu	Ser	Phe	Met	Leu 15	Leu
15	Phe	Ser	Leu	Trp 20	Arg	Gln	Ser	Суз	Arg 25	Arg	Arg	Lys	Leu	Pro 30	Pro	Gly
	Pro	Thr	Pro 35	Leu	Pro	Ile	Ile	Gly 40	Asn	Met	Leu	Gln	Ile 45	Asp	Val	Lys
20	Asp	Ile 50	Cys	Lys	Ser	Phe	Thr 55	Asn	Phe	Ser	Lys	Val 60	Tyr	Gly	Pro	Val
	Phe 65	Thr	Val	Tyr	Phe	Gly 70	Met	Asn	Pro	Ile	Val 75	Val	Phe	His	Gly	Tyr 80
25	Glu	Ala	Val	Lys	Glu 85	Ala	Leu	Ile	Asp	Asn 90	Gly	Glu	Glu	Phe	Ser 95	Gly
	Arg	Gly	Asn	Ser 100	Pro	Ile	Ser	Gln	Arg 105	Ile	Thr	Lys	Gly	Leu 110	Gly	Ile
30	Ile	Ser	Ser 115	Asn	Gly	Lys	Arg	Trp 120	Lys	Glu	Ile	Arg	Arg 125	Phe	Ser	Leu
	Thr	Thr 130	Leu	Arg	Asn	Phe	Gly 135	Met	Gly	Lys	Lys	Ser 140	Ile	Glu	Asp	Arg
35	Val 145	Gln	Glu	Glu	Ala	His 150	Cys	Leu	Val	Glu	Glu 155	Leu	Arg	Lys	Thr	Lys 160
	Ala	Ser	Pro	Cys	Asp 165	Pro	Thr	Phe	Ile	Leu 170	Gly	Cys	Ala	Pro	Cys 175	Asn
40	Val	Ile	Cys	Ser 180	Val	Val	Phe	Gln	Lys 185	Arg	Phe	Asp	Tyr	Lys 190	Asp	Gln
<b>4</b> 5	Asn	Phe	Leu 195	Thr	Leu	Met	Lys	Arg 200	Phe	Asn	Glu	Asn	Phe 205	Arg	Ile	Leu
.•	Asn	Ser 210	Pro	Trp	Ile	Gln	Val 215	Cys	Asn	Asn	Phe	Pro 220	Leu	Leu	Ile	Asp
50	Cys 225	Phe	Pro	Gly	Thr	His 230	Asn	Lys	Val	Leu	Lys 235	Asn	Val	Ala	Leu	Thr 240

	Arg	Ser	Tyr	Ile	Arg 245	Glu	Lys	Val	Lys	Glu 250	His	Gln	Ala	Ser	Leu 255	Asp
5	Val	Asn	Asn	Pro 260	Arg	Asp	Phe	Ile	Asp 265	Cys	Phe	Leu	Ile	Lys 270	Met	Glu
	Gln	Glu	Lys 275	Asp	Asn	Gln	Lys	Ser 280	Glu	Phe	Asn	Ile	Glu 285	Asn	Leu	Val
10	Gly	Thr 290	Val	Ala	Asp	Leu	Phe 295	Val	Ala	Gly	Thr	Glu 300	Thr	Thr	Ser	Thr
	Thr 305	Leu	Arg	Tyr	Gly	Leu 310	Leu	Leu	Leu	Leu	Lys 315	His	Pro	Glu	Val	Thr 320
15	Ala	Lys	Val	Gln	Glu 325	Glu	Ile	Asp	His	Val 330	Ile	Gly	Arg	His	Arg 335	Ser
20	Pro	Cys	Met	Gln 340	Asp	Arg	Ser	His	Met 345	Pro	Tyr	Thr	Asp	Ala 350	Val	Val
	His	Glu	Ile 355	Gln	Arg	Tyr	Ser	Asp 360	Leu	Val	Pro	Thr	Gly 365	Val	Pro	His
25	Ala	<b>V</b> al 370	Thr	Thr	Asp	Thr	Lys 375	Phe	Arg	Asn	Tyr	Leu 380	Ile	Pro	Lys	Gly
	Thr 385	Thr	Ile	Met	Ala	Leu 390	Leu	Thr	Ser	Val	Leu 395	His	Asp	Asp	Arg	Glu 400
30	Phe	Pro	Asn	Pro	Asn 405	Ile	Phe	Asp	Pro	Gly 410	His	Phe	Leu	Asp	Lys 415	Asn
	Gly	Asn	Phe	Lys 420	Lys	Ser	Asp	Tyr	Phe 425	Met	Pro	Phe	Ser	Ala 430	Gly	Lys
35	Arg	Ile	Cys 435	Ala	Gly	Glu	Gly	Leu 440	Ala	Arg	Met	Glu	Leu 445	Phe	Leu	Phe
	Leu	Thr 450	Thr	Ile	Leu	Gln	<b>As</b> n <b>4</b> 55	Phe	Asn	Leu	Lys	Ser 460	Val	Asp	Asp	Leu
40	Lys 465	Asn	Leu	Asn	Thr	Thr 470	Ala	Val	Thr	Lys	Gly 475	Ile	Val	Ser	Leu	Pro 480
<b>4</b> 5	Pro	Ser	Tyr	Gln	Ile 485	Cys	Phe	Ile	Pro	Val 490						
~	(2)					SEQ										
50		(i)	( I ( E	A) LE B) TY C) ST	ENGTH	HARAC H: 14 nucl DEDNE DGY:	173 k Leic ESS:	ase acio doub	pair I	s						

(ix) FEATURE:
(A) NAME/KEY: CDS
(B) LOCATION: 1..1470

5		(xi)	SEC	QUEN	CE DI	ESCR	ITTI	) <b>n</b> : 5	SEQ :	D NO	): 25	ō:						
	ATG Met 1	GAA Glu	CCT Pro	TTT Phe	GTG Val 5	GTC Val	CTG Leu	GTG <b>Va</b> l	CTG Leu	TGT Cys 10	CTC Leu	TCT Ser	TTT Phe	ATG Met	CTT Leu 15	CTC Leu		48
10	TTT Phe	TCA Ser	CTC Leu	TGG Trp 20	AGA Arg	C <b>A</b> G Gln	AGC Ser	TGT Cys	AGG Arg 25	AGA Arg	AGG Arg	AAG Lys	CTC Leu	CCT Pro 30	CCT Pro	GGC Gly		96
15				CTT Leu													:	144
	GAC Asp	ATC Ile 50	TGC Cys	AAA Lys	TCT Ser	TTC Phe	ACC Thr 55	<b>A</b> AT <b>A</b> sn	TTC Phe	TCA Ser	AAA Lys	GTC Val 60	TAT Tyr	GGT Gly	CCT Pro	GTG Val	;	192
20				TAT Tyr													:	240
25	GTG Val	GCA Ala	GTG Val	AAG Lys	GAA Glu 85	GCC Ala	CTG Leu	ATT Ile	GAT Asp	AAT Asn 90	GGA Gly	GAG Glu	GAG Glu	TTT Phe	TCT Ser 95	GGA Gly	:	288
				TCC Ser 100													:	336
30	ATT Ile	TCC Ser	AGC Ser 115	AAT Asn	GGA Gly	AAG Lys	AGA Arg	TGG Trp 120	AAG Lys	GAG Glu	ATC Ile	CGG Arg	CGT Arg 125	TTC Phe	TCC Ser	CTC Leu	<u>-</u>	384
35	ACA Thr	ACC Thr 130	TTG Leu	CGG Arg	AAT Asn	TTT Phe	GGG Gly 135	ATG Met	GGG Gly	AAG Lys	AAG Lys	AGC Ser 140	ATT Ile	GAG Glu	GAC Asp	CGT Arg	•	432
	GTT Val 145	CAA Gln	GAG Glu	GAA Glu	GCT Ala	CAC His 150	TGC Cys	CTT Leu	GTG Val	GAG Glu	GAG Glu 155	TTG Leu	AGA Arg	AAA Lys	ACC Thr	AAG Lys 160	,	480
40	GCT Ala	TCA Ser	CCC Pro	TGT Cys	GAT Asp 165	CCC Pro	ACT Thr	TTC Phe	ATC Ile	CTG Leu 170	GGC Gly	TGT Cys	GCT Ala	CCC Pro	TGC Cys 175	AAT Asn	!	528
<b>4</b> 5				TCC Ser 180													!	576
				ACC Thr													1	624

50

										Leu		GAT Asp	672
5				CAC His 230					Asn			ACA Thr 240	720
10				GAG Glu									768
				GAC Asp									816
15				CAA Gln									864
20				CTA Leu									912
				CTC Leu 310									960
25				GAG Glu									1008
30				AGG Arg									1056
	_			TAC Tyr									1104
35	_	_		ACT Thr									1152
40				TTA Leu 390									1200
	_		Pro	ATC Ile		Asp	Gly	His			Asp		1248
<b>4</b> 5				AGT Ser									1296
50				GAA Glu	Gly								1344

	CTA Leu	ACC Thr 450	ACA Thr	ATT Ile	TTA Leu	CAG Gln	AAC Asn 455	TTT Phe	AAC Asn	CTG Leu	AAA Lys	TCT Ser 460	GTT Val	GAT Asp	GAT Asp	TTA Leu	1392
5	AAG Lys 465	AAC Asn	CTC Leu	AAT Asn	ACT Thr	ACT Thr 470	GCA Ala	GTT Val	ACC Thr	AAA Lys	GGG Gly 475	ATT Ile	GTT Val	TCT Ser	CTG Leu	CCA Pro 480	1440
10		TCA Ser									TGA						1473
15	(2) INFORMATION FOR SEQ ID NO: 26:  (i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 490 amino acids  (B) TYPE: amino acid  (D) TOPOLOGY: linear																
20	(ii) MOLECULE TYPE: protein  (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 26:																
	Met 1	Glu	Pro	Phe	Val 5	Val	Leu	Val	Leu	Cys 10	Leu	Ser	Phe	Met	Leu 15	Leu	
25	Phe	Ser	Leu	Trp 20	Arg	Gln	Ser	Cys	Arg 25	Arg	Arg	Lys	Leu	Pro 30	Pro	Gly	
	Pro	Thr	Pro 35	Leu	Pro	Ile	Ile	Gly 40	Asn	Met	Leu	Gln	Ile 45	Asp	Val	Lys	
30	Asp	Ile 50	Cys	Г <b>ү</b> в	Ser	Phe	Thr 55	Asn	Phe	Ser	Lys	Val 60	Tyr	Gly	Pro	Val	
	Phe 65	Thr	Val	Tyr	Phe	Gly 70	Met	Asn	Pro	Ile	<b>Val</b> 75	Val	Phe	His	Gly	Tyr 80	
35	Val	Ala	Val	Lys	Glu 85	Ala	Leu	Ile	Asp	Asn 90	Gly	Glu	Glu	Phe	Ser 95	Gly	
	Arg	Gly	Asn	Ser 100	Pro	Ile	Ser	Gln	Arg 105	Ile	Thr	Lys	Gly	Leu 110	Gly	Ile	
<b>4</b> 0	Ile	Ser	Ser 115	Asn	Gly	Lys	Arg	Trp 120	Lys	Glu	Ile	Arg	Arg 125	Phe	Ser	Leu	
	Thr	Thr 130	Leu	Arg	Asn	Phe	Gly 135		Gly	Lys	Lys	Ser 140	Ile	Glu	Asp	Arg	
<b>4</b> 5	Val 145	Gln	Glu	Glu	Ala	His 150		Leu	Val	Glu	Glu 155	Leu	Arg	Lys	Thr	Lys 160	
	Ala	Ser	Pro	Cys	Asp 165		Thr	Phe	Ile	Leu 170		Cys	Ala	Pro	Cys 175	Asn	

	Val	Ile	Cys	Ser 180	Val	Val	Phe	Gln	Lys 185	Arg	Phe	Asp	Tyr	Lys 190	Asp	Gln
5	Asn	Phe	Leu 195	Thr	Leu	Met	Lys	Arg 200	Phe	Asn	Glu	Asn	Phe 205	Arg	Ile	Leu
	Asn	Ser 210	Pro	Trp	Ile	Gln	Val 215	Cys	Asn	Asn	Phe	Pro 220	Leu	Leu	Ile	Asp
10	Cys 225	Phe	Pro	Gly	Thr	His 230	Asn	Lys	Val	Leu	Lys 235	Asn	Val	Ala	Leu	Thr 240
	Arg	Ser	Tyr	Ile	Arg 245	Glu	Lys	Val	Lys	Glu 250	His	Gln	Ala	Ser	Leu 255	Asp
15	Val	Asn	Asn	Pro 260	Arg	Asp	Phe	Ile	Asp 265	Cys	Phe	Leu	Ile	Lys 270	Met	Glu
	Gln	Glu	Lys 275	Asp	Asn	Gln	Lys	Ser 280	Glu	Phe	Asn	Ile	Glu 285	Asn	Leu	Val
20	Gly	Thr 290	Val	Ala	Asp	Leu	Phe 295	Val	Ala	Gly	Thr	Glu 300	Thr	Thr	Ser	Thr
	Thr 305	Leu	Arg	Tyr	Gly	Leu 310	Leu	Leu	Leu	Leu	<b>Lys</b> 315	His	Pro	Glu	Val	Thr 320
25	Ala	Lys	Val	Gln	Glu 325	Glu	Ile	Asp	His	Val 330	Ile	Gly	Arg	His	Arg 335	Ser
	Pro	Cys	Met	Gln 340	Asp	Arg	Ser	His	Met 345	Pro	Tyr	Thr	Asp	Ala 350	Val	Val
30	His	Glu	Ile 355	Gln	Arg	Tyr	Ser	Asp 360	Leu	Val	Pro	Thr	Gly 365	Val	Pro	His
	Ala	Val 370	Thr	Thr	Asp	Thr	Lys 375	Phe	Arg	Asn	Tyr	Leu 380	Ile	Pro	Lys	Gly
35	Thr 385	Thr	Ile	Met	Ala	Leu 390	Leu	Thr	Ser	Val	Leu 395	His	Asp	Asp	Arg	Glu 400
	Phe	Pro	Asn	Pro	Asn 405	Ile	Phe	Asp	Pro	Gly 410	His	Phe	Leu	Asp	Lys 415	Asn
40	Gly	Asn	Phe	Lys 420	Lys	Ser	Asp	Tyr	Phe 425	Met	Pro	Phe	Ser	Ala 430	Gly	Lys
	Arg	Ile	Cys 435	Ala	Gly	Glu	Gly	Leu 440	Ala	Arg	Met	Glu	Leu 445	Phe	Leu	Phe
<b>4</b> 5	Leu	Thr 450	Thr	Ile	Leu	Gln	<b>A</b> sn 455	Phe	Asn	Leu	Lys	Ser 460	Val	Asp	Asp	Leu
	Lys 465	Asn	Leu	Asn	Thr	Thr 470	Ala	Val	Thr	Lys	Gly 475	Ile	Val	Ser	Leu	Pro 480
50	Pro	Ser	Tyr	Gln	Ile 485	Cys	Phe	Ile	Pro	Val 490						

	(2)	INF	ORMA'	TION	FOR	SEQ	ID I	NO: 3	27:								
5		(i)	() () ()	QUENCA) LI B) T C) S D) T	ENGTI YPE : FRANI	H: 14 nucl DEDNI	473 l leic ESS:	base acid doul	pai:	rs							
10			() ()	ATURI A) NA B) L	AME/I	ION:	1		ano :	TD M	<b>1.</b> 21	7.					
15	ATG Met 1	GAT	CCA	QUEN( GCT Ala	GTG	GCT	CTG	GTG	CTC	TGT	CTC	TCC	TGT Cys	TTG Leu	TTT Phe 15	CTC Leu	4.8
	CTT Leu	TCA Ser	CTC Leu	TGG Trp 20	AGG Arg	CAG Gln	AGC Ser	TCT Ser	GGA Gly 25	AGA Arg	GGG Gly	AGG Arg	CTC Leu	CCG Pro 30	TCT Ser	GGC Gly	96
20	CCC Pro	ACT Thr	CCT Pro 35	CTC Leu	CCG Pro	ATT Ile	ATT Ile	GGA Gly 40	AAT Asn	ATC Ile	CTG Leu	CAG Gln	TTA Leu 45	GAT Asp	GTT Val	AAG Lys	144
25				AAA Lys													192
				TAT Tyr													240
30	GAA Glu	GCA Ala	GTG <b>V</b> al	AAG Lys	GAG Glu 85	GCC Ala	CTG Leu	ATT Ile	GAT Asp	CAT His 90	GGA Gly	GAG Glu	GAG Glu	TTT Phe	TCT Ser 95	GGA Gly	288
35	AGA Arg	GGA Gly	AGT Ser	TTT Phe 100	CCA Pro	GTG Val	GCT Ala	GAA Glu	AAA Lys 105	GTT Val	AAC Asn	AAA Lys	GGA Gly	CTT Leu 110	GGA Gly	ATC Ile	336
	CTT Leu	TTC Phe	AGC Ser 115	AAT Asn	GGA Gly	AAG Lys	AGA Arg	TGG Trp 120	AAG Lys	GAG Glu	ATC Ile	CGG Arg	CGT Arg 125	TTC Phe	TGC Cys	CTC Leu	384
<b>4</b> 0				CGG Arg									_	_			432
<b>4</b> 5				GAA Glu													480
50																	

	GCC Ala	TCA Ser	CCC Pro	TGT Cys	GAT Asp 165	CCC Pro	ACT Thr	TTC Phe	ATC Ile	CTG Leu 170	GGC Gly	TGT Cys	GCT Ala	CCC Pro	TGC Cys 175	AAT Asn		528
5	GTG Val	ATC Ile	TGC Cys	TCT Ser 180	GTT Val	ATT Ile	TTC Phe	CAT His	GAT Asp 185	CGA Arg	TTT Phe	GAT Asp	TAT Tyr	AAA Lys 190	GAT Asp	CAG Gln		576
10						ATG <b>M</b> et												624
						CAG Gln												672
15						CAT His 230												720
20						GAG Glu												768
	ATG Met	AAC Asn	AGT Ser	GCT Ala 260	CGG Arg	GAC Asp	TTT Phe	ATT Ile	GAT Asp 265	TGT Cys	TTC Phe	CTG Leu	ATC Ile	AAA Lys 270	ATG Met	GAA Glu		816
25						CAA Gln												864
30						ATG Met												912
	ACT Thr 305	CTG Leu	AGA Arg	TAT Tyr	GGA Gly	CTC Leu 310	CTG Leu	CTC Leu	CTG Leu	CTG Leu	AAG Lys 315	TAC Tyr	CCA Pro	GAG Glu	GTC Val	ACA Thr 320		960
35						GAG Glu										AGC Ser	-	1008
40						AGG Arg											j	1056
						TAC Tyr											-	1104
<b>4</b> 5						GTT Val											Ī	1152
50						TCC Ser 390											3	1200

	TTC Phe	CCC Pro	AAC Asn	CCA Pro	GAG Glu 405	ATG Met	TTT Phe	GAC Asp	CCT Pro	GGC Gly 410	CAC His	TTT Phe	CTG Leu	GAT Asp	AAG Lys 415	AGT Ser	:	1248
5	GGC Gly	AAC Asn	TTT Phe	AAG Lys 420	AAA Lys	AGT Ser	GAC Asp	TAC Tyr	TTC Phe 425	ATG Met	CCT Pro	TTC Phe	TCA Ser	GCA Ala 430	GGA Gly	AAA Lys	:	1296
10	CGG Arg	ATG Met	TGT Cys 435	ATG Met	GGA Gly	GAG Glu	GGC Gly	CTG Leu 440	GCC Ala	CGC Arg	ATG Met	GAG Glu	CTG Leu 445	TTT Phe	TTA Leu	TTC Phe	:	1344
	CTG Leu	ACC Thr 450	ACC Thr	ATT Ile	TTG Leu	CAG Gln	AAC Asn 455	TTT Phe	AAC Asn	CTG Leu	AAA Lys	TCT Ser 460	CAG Gln	GTT Val	GAC Asp	CCA Pro	:	1392
15	AAG Lys 465	GAT Asp	ATT Ile	GAC Asp	ATC Ile	ACC Thr 470	CCC Pro	ATT Ile	GCC Ala	AAT Asn	GCA Ala 475	TTT Phe	GGT Gly	CGT Arg	GTG Val	CCA Pro 480		1440
20	CCC Pro	TTG Leu	TAC Tyr	CAG Gln	CTC Leu 485	TGC Cys	TTC Phe	ATT Ile	CCT Pro	GTC Val 490	TGA						:	1473
	(2)																	
25	CCC TTG TAC CAG CTC TGC TTC ATT CCT GTC TGA Pro Leu Tyr Gln Leu Cys Phe Ile Pro Val 485  (2) INFORMATION FOR SEQ ID NO: 28:  (i) SEQUENCE CHARACTERISTICS: (A) LENGTH: 490 amino acids (B) TyPE: amino acid (D) TOPOLOGY: linear  (ii) MOLECULE TYPE: protein (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 28:																	
									SEQ I	ED <b>N</b> C	D: 28	3:						
30	Met 1	Asp	Pro	Ala	Val 5	Ala	Leu	Val	Leu	Cys 10	Leu	Ser	Cys	Leu	Phe 15	Leu		
35	Leu	Ser	Leu	Trp 20	Arg	Gln	Ser	Ser	Gly 25	Arg	Gly	Arg	Leu	Pro 30	Ser	Gly '		
00	Pro	Thr	Pro 35	Leu	Pro	Ile	Ile	Gly 40	Asn	Ile	Leu	Gln	Leu 45	Asp	Val	Lys		
40	Asp	<b>M</b> et 50	Ser	Lys	Ser	Leu	Thr 55	Asn	Phe	Ser	Lys	Val 60	Tyr	Gly	Pro	Val		
	Phe 65	Thr	Val	Tyr	Phe	Gly 70	Leu	Lys	Pro	Ile	<b>Val</b> 75	Val	Leu	His	Gly	Tyr 80		
45	Glu	Ala	Val	Lys	Glu 85	Ala	Leu	Ile	Asp	His 90	Gly	Glu	Glu	Phe	Ser 95	Gly		
~	Arg	Gly	Ser	Phe 100	Pro	Val	Ala	Glu	<b>Lys</b> 105	Val	Asn	Lys	Gly	Leu 110	Gly	Ile		

	Leu	Phe	Ser 115	Asn	Gly	Lys	Arg	Trp 120	Lys	Glu	Ile	Arg	<b>A</b> rg 125	Phe	Cys	Leu
5	Met	Thr 130	Leu	Arg	Asn	Phe	Gly 135	Met	Gly	Lys	Arg	Ser 140	Ile	Glu	Asp	Arg
	Val 145	Gln	Glu	Glu	Ala	Arg 150	Cys	Leu	Val	Glu	Glu 155	Leu	Arg	Lys	Thr	Asn 160
10	Ala	Ser	Pro	Cys	Asp 165	Pro	Thr	Phe	Ile	Leu 170	Gly	Cys	Ala	Pro	Cys 175	Asn
	Val	Ile	Сув	Ser 180	Val	Ile	Phe	His	Asp 185	Arg	Phe	Asp	Tyr	Lys 190	Asp	Gln
15	Arg	Phe	Leu 195	Asn	Leu	Met	Glu	Lys 200	Phe	Asn	Glu	Asn	Leu 205	Arg	Ile	Leu
	Ser	Ser 210	Pro	Trp	Ile	Gln	Val 215	Суѕ	Asn	Asn	Phe	Pro 220	Ala	Leu	Ile	Asp
20	Tyr 225	Leu	Pro	Gly	Ser	His 230	Asn	Lys	Ile	Ala	Glu 235	Asn	Phe	Ala	Tyr	Ile 240
	Lys	Ser	Tyr	Val	Leu 245	Glu	Arg	Ile	Lys	Glu 250	His	Gln	Glu	Ser	Leu 255	Asp
25	Met	Asn	Ser	Ala 260	Arg	Asp	Phe	Ile	Asp 265	Cys	Phe	Leu	Ile	Lys 270	Met	Glu
	Gln	Glu	Lys 275	His	Asn	Gln	Gln	Ser 280	Glu	Phe	Thr	Val	Glu 285	Ser	Leu	Ile
30	Ala	Thr 290	Val	Thr	Asp	Met	Phe 295	Gly	Ala	Gly	Thr	Glu 300	Thr	Thr	Ser	Thr
	Thr 305	Leu	Arg	Tyr	Gly	Leu 310	Leu	Leu	Leu	Leu	Lys 315	Tyr	Pro	Glu	Val	Thr 320
35	Ala	Lys	Val	Gln	Glu 325	Glu	Ile	Glu	Cys	<b>Val</b> 330	Val	Gly	Arg	Asn	Arg 335	Ser
	Pro	Cys	Met	Gln 340	Asp	Arg	Ser	His	Met 345	Pro	Tyr	Thr	Asp	Ala 350	Val	Val
40	His	Glu	Ile 355	Gln	Arg	Туr	Ile	Asp 360	Leu	Leu	Pro	Thr	Asn 365	Leu	Pro	His
	Ala	Val 370	Thr	Cys	Asp	Val	Lys 375	Phe	Lys	Asn	Tyr	Leu 380	Ile	Pro	Lys	Gly
<b>4</b> 5	Thr 385	Thr	Ile	Ile	Thr	Ser 390	Leu	Thr	Ser	Val	Leu 395	His	Asn	qaA	Lys	Glu 400
	Phe	Pro	Asn	Pro	Glu 405	Met	Phe	Asp	Pro	Gly 410	His	Phe	Leu	Asp	Lys 415	Ser
50	Gly	Asn	Phe	Lys 420	Lys	Ser	Asp	Tyr	Phe 425	Met	Pro	Phe	Ser	Ala 430	Gly	Lys

	Arg	Met	Cys 435	Met	Gly	Glu	Gly	Leu 440	Ala	Arg	Met	Glu	Leu 445	Phe	Leu	Phe	
5	Leu	Thr 450	Thr	Ile	Leu	Gln	Asn 455	Phe	Asn	Leu	Lys	Ser 460	Gln	Val	Asp	Pro	
	Lys 465	Asp	Ile	Asp	Ile	Thr 470	Pro	Ile	Ala	Asn	Ala 475	Phe	Gly	Arg	Val	Pro 480	
10	Pro	Leu	Tyr	Gln	Leu 485	Сув	Phe	Ile	Pro	Val 490							
	(2)	INFO	ORMA'	rion	FOR	SEQ	ID N	10: 2	29:								
15		(i)	() (E	QUENCA) LE B) TY C) ST O) TO	ENGTI PE: PRANI	H: 14 nucl	173 h leic ESS:	ase acid doub	pain l	rs							
20		(ix)	(2	ATURI A) NA 3) LO	AME/I			470									
		( <b>x</b> i)	SEC	QUENC	CE DE	ESCRI	PTIC	ON: S	SEQ I	D NO	): 29	9 :					
25				TTT Phe													4.8
30				TGG Trp 20													96
				CTC Leu								_			_		144
35				AAA Lys													192
_				TAT Tyr													240
40				AAG Lys													288
<b>4</b> 5				TTC Phe 100												_	336
50																	

	GTT Val	TTC Phe	AGC Ser 115	AAT Asn	GGA Gly	AAG Lys	AGA Arg	TGG Trp 120	AAG Lys	GAG Glu	ATC Ile	CGG <b>A</b> rg	CGT Arg 125	TTC Phe	TCC Ser	CTC Leu	384
5	ATG Met	ACG Thr 130	CTG Leu	CGG <b>A</b> rg	AAT Asn	TTT Phe	GGG Gly 135	ATG Met	GGG Gly	AAG Lys	AGG Arg	AGC Ser 140	ATT Ile	GAG Glu	GAC Asp	CGT Arg	432
10	GTT Val 145	CAA Gln	GAG Glu	G <b>AA</b> Glu	GCC <b>A</b> la	CGC Arg 150	TGC Cys	CTT Leu	GTG Val	GAG Glu	GAG Glu 155	TTG Leu	AGA Arg	AAA Lys	ACC Thr	AAG Lys 160	480
10	GCT Ala	TCA Ser	CCC Pro	TGT Cys	GAT Asp 165	CCC Pro	ACT Thr	TTC Phe	ATC Ile	CTG Leu 170	GGC Gly	TGT Cys	GCT Ala	CCC Pro	TGC Cys 175	AAT Asn	528
15	GTG Val	ATC Ile	TGC Cys	TCC Ser 180	ATT Ile	ATT Ile	TTC Phe	CAG Gln	AAA Lys 185	CGT Arg	TTC Phe	GAT Asp	TAT Tyr	AAA Lys 190	GAT Asp	CAG Gln	576
												AAC Asn					624
20	AGC Ser	ACC Thr 210	CCC Pro	TGG Trp	ATC Ile	CAG Gln	ATA Ile 215	TGC Cys	AAT Asn	AAT Asn	TTT Phe	CCC Pro 220	ACT Thr	ATC Ile	ATT Ile	GAT Asp	672
25												AAC Asn					720
												CAA Gln					768
30												CTG Leu					816
35												ATT Ile				GTA Val	864
												GAG Glu 300					912
40												CAC His					960
<b>4</b> 5												GGC Gly					1008
												ACA Thr					1056

		GAG Glu															1104
5	GCA Ala	GTG Val 370	ACC Thr	TGT Cys	GAC Asp	GTT Val	AAA Lys 375	TTC Phe	AGA Arg	AAC Asn	TAC Tyr	CTC Leu 380	ATT Ile	CCC Pro	AAG Lys	GGC Gly	1152
10		ACC Thr															1200
		CCC Pro															1248
15		AAT Asn															1296
20		ATT Ile															1344
20		ACC Thr 450															1392
25		GAC Asp															1440
30		TTC Phe								_	TGA						1473
	(2)	INFO	RMAT	ON	FOR	SEQ	ID N	<b>10:</b> 3	0:								
35		(	(A (B	L) LE	NGTH	CHAF I: 49 amir GY:	00 am	nino cid								,	
		(ii)	MOL	ECUI	E TY	PE:	prot	ein									
<b>4</b> 0						SCRI			•						<b>.</b>		
	Met 1	Asp	Pro	Phe	Va 1 5	Val	Leu	Val	Leu	Cys 10	Leu	Ser	Cys	Leu	ьеи 15	Leu	
	Leu	Ser	Leu	Trp 20	Arg	Gln	Ser	Ser	Gly 25	Arg	Gly	Lys	Leu	Pro 30	Pro	Gly	
<b>4</b> 5	Pro	Thr	Pro 35	Leu	Pro	Val	Ile	Gly 40	Asn	Ile	Leu	Gln	Ile 45	Asp	Ile	Lys	

	Asp	Val 50	Ser	Lys	Ser	Leu	Thr 55	Asn	Leu	Ser	Lys	Ile 60	Tyr	Gly	Pro	Val
5	Phe 65	Thr	Leu	Tyr	Phe	Gly 70	Leu	Glu	Arg	Met	Val 75	Val	Leu	His	Gly	Tyr 80
	Glu	Val	Val	Lys	Glu 85	Ala	Leu	Ile	Asp	Leu 90	Gly	Glu	Glu	Phe	Ser 95	Gly
10	Arg	Gly	His	Phe 100	Pro	Leu	Ala	Glu	Arg 105	Ala	Asn	Arg	Gly	Phe 110	Gly	Ile
		Phe	115					120					125			
15		Thr 130					135					140				
	145	Gln				150					155					160
20		Ser			165					170					175	
		Ile		180					185					190		
25		Phe	195					200					205			
		Thr 210					215					220				
30	225	Phe				230					235					240
		Ser			245					250					255	
35		Asn		260					265					270		
40	-	Glu	275					280					285			
40		Thr 290					295					300				
45	305					310					315					Thr 320
<b>4</b> 5		Lys			325					330					335	
50	Pro	Cys	Met	Gln 340	Asp	Arg	Gly	His	Met 345	Pro	Tyr	Thr	Asp	Ala 350	Val	Val
<i>5</i> 0	His	Glu	<b>Va</b> l		Arg	Tyr	Ile	Asp 360	Leu	Ile	Pro	Thr	Ser 365	Leu	Pro	His

	Ala	Val 370	Thr	Cys	Asp	Val	Lys 375	Phe	Arg	Asn	Tyr	Leu 380	Ile	Pro	Lys	Gly	
5	Thr 385	Thr	Ile	Leu	Thr	Ser 390	Leu	Thr	Ser	Val	Leu 395	His	Asp	Asn	Lys	Glu 400	
	Phe	Pro	Asn	Pro	Glu 405	Met	Phe	Asp	Pro	Arg 410	His	Phe	Leu	Asp	Glu 415	Gly	
10	Gly	Asn	Phe	Lys 420	Lys	Ser	Asn	Tyr	Phe 425	Met	Pro	Phe	Ser	Ala 430	Gly	Lys	
	Arg	Ile	Cys 435	Val	Gly	Glu	Gly	Leu 440	Ala	Arg	Met	Glu	Leu 445	Phe	Leu	Phe	
15	Leu	Thr 450	Phe	Ile	Leu	Gln	Asn 455	Phe	Asn	Leu	Lys	Ser 460	Leu	Ile	Asp	Pro	
	Lys 465	Asp	Leu	Asp	Thr	Thr 470	Pro	Val	Val	Asn	Gly 475	Phe	Ala	Ser	Val	Pro 480	
20	Pro	Phe	Tyr	Gln	Leu 485	Cys	Phe	Ile	Pro	Val 490							
	(2)																
<b>2</b> 5	Pro Phe Tyr Gln Leu Cys Phe Ile Pro Val																
30		(ix)	(1	A) NA	ME/I			1491									
		(xi)	SEÇ	QUENC	CE DE	ESCRI	PTIC	ON: 5	SEQ :	D NO	): 31	l:					
35				GAA Glu													48
40				GTG Val 20													96
		Pro	Pro	GGC Gly	Pro	Leu	Pro	Leu	Pro	Gly	Leu	Gly	Asn				144
<b>4</b> 5				CAG Gln													192
50																	

	TTC Phe 65	GGG Gly	GAC Asp	GTG Val	TTC Phe	AGC Ser 70	CTG Leu	CAG Gln	CTG Leu	GCC Ala	TGG Trp 75	ACG Thr	CCG Pro	GTG Val	GTC Val	GTG Val 80	:	240
5	CTC Leu	AAT Asn	GGG Gly	CTG Leu	GCG Ala 85	GCC Ala	GTG Val	CGC <b>A</b> rg	GAG Glu	GCG Ala 90	CTG Leu	GTG Val	ACC Thr	CAC His	GGC Gly 95	GAG Glu	;	288
10	GAC Asp	ACC Thr	GCC Ala	GAC Asp 100	CGC <b>A</b> rg	CCG Pro	CCT Pro	GTG Val	CCC Pro 105	ATC Ile	ACC Thr	CAG Gln	ATC Ile	CTG Leu 110	GGT Gly	TTC Phe		336
	GGG Gly	CCG Pro	CGT Arg 115	TCC Ser	CAA Gln	GGG Gly	GTG Val	TTC Phe 120	CTG Leu	GCG Ala	CGC Arg	TAT Tyr	GGG Gly 125	CCC Pro	GCG Ala	TGG Trp	:	384
15	CGC Arg	GAG Glu 130	CAG Gln	AGG Arg	CGC Arg	TTC Phe	TCC Ser 135	GTC Val	TCC Ser	ACC Thr	TTG Leu	CGC Arg 140	AAC Asn	TTG Leu	GGC Gly	CTG Leu	•	432
20	GGC Gly 145	<b>AA</b> G Lys	AAG Lys	TCG Ser	CTG Leu	GAG Glu 150	CAG Gln	TGG Trp	GTG Val	ACC Thr	GAG Glu 155	GAG Glu	GCC Ala	GCC Ala	TGC Cys	CTT Leu 160	•	480
	TGT Cys	GCC Ala	GCC Ala	TTC Phe	GCC Ala 165	AAC Asn	CAC His	TCC Ser	GGA Gly	CGC Arg 170	CCC Pro	TTT Phe	CGC <b>A</b> rg	CCC Pro	AAC Asn 175	GGT Gly	!	528
25	CTC Leu	TTG Leu	GAC Asp	AAA Lys 180	GCC Ala	GTG Val	AGC Ser	AAC Asn	GTG Val 185	ATC Ile	GCC Ala	TCC Ser	CTC Leu	ACC Thr 190	TGC Cys	GGG Gly	!	576
30	CGC Arg	CGC Arg	TTC Phe 195	GAA Glu	TAC Tyr	GAC Asp	GAC Asp	CCT Pro 200	CGC <b>Ar</b> g	TTC Phe	CTC Leu	AGG Arg	CTG Leu 205	CTG Leu	GAC Asp	CTA Leu	(	624
	Ala	Gln 210	Glu	Gly	Leu	Lys	Glu 215	Glu	Ser	Gly		Leu 220	Arg	Glu	Val	Leu	1	672
35	Asn 225	Ala	Val	Pro	Val	Leu 230	Leu	His	Ile	Pro	GCG Ala 235	Leu	Ala	Gly	Lys	Val 240		720
40	Leu	Arg	Phe	Gln	Lys 245	Ala	Phe	Leu	Thr	Gln 250	CTG Leu	Asp	Glu	Leu	Leu 255	Thr	,	768
	GAG Glu	CAC His	AGG Arg	ATG Met 260	Thr	TGG Trp	Asp	Pro	GCC Ala 265	Gln	CCC Pro	CCC Pro	Arg	Asp	Leu	ACT Thr	1	816
<b>4</b> 5	GAG Glu	GCC Ala	TTC Phe 275	CTG Leu	GCA Ala	GAG Glu	ATG Met	GAG Glu 280	AAG Lys	GCC Ala	AAG Lys	GGG Gly	AAC Asn 285	CCT Pro	GAG Glu	AGC Ser	i	864
50	AGC Ser	TTC Phe 290	AAT Asn	GAT <b>A</b> sp	GAG Glu	AAC Asn	CTG Leu 295	TGC Cys	ATA Ile	GTG Val	GTG Val	GCT Ala 300	GAC Asp	CTG Leu	TTC Phe	TCT Ser		912

	GCC Ala 305	GGG Gly	ATG Met	GTG Val	ACC Thr	ACC Thr 310	TCG Ser	ACC Thr	ACG Thr	CTG Leu	GCC Ala 315	TGG Trp	GGC Gly	CTC Leu	CTG Leu	CTC Leu 320	960
5	ATG Met	ATC Ile	CTA Leu	CAT His	CCG Pro 325	GAT Asp	GTG Val	CAG Gln	CGC Arg	CGT Arg 330	GTC <b>V</b> al	CAA Gln	CAG Gln	GAG Glu	ATC Ile 335	GAC Asp	1008
10	GAC Asp	GTG Val	ATA Ile	GGG Gly 340	CAG Gln	GTG Val	CGG Arg	CGA Arg	CCA Pro 345	GAG Glu	ATG Met	GGT Gly	GAC Asp	CAG Gln 350	GCT Ala	CAC His	1056
	ATG Met	CCC Pro	TAC Tyr 355	ACC Thr	ACT Thr	GCC Ala	GTG Val	ATT Ile 360	CAT His	GAG Glu	GTG Val	CAG Gln	CGC Arg 365	TTT Phe	GGG Gly	GAC Asp	1104
15	ATC Ile	GTC Val 370	CCC Pro	CTG Leu	GGT Gly	GTG Val	ACC Thr 375	CAT His	ATG Met	ACA Thr	TCC Ser	CGT Arg 380	GAC Asp	ATC Ile	GAA Glu	GTA Val	1152
20	CAG Gln 385	GGC Gly	TTC Phe	CGC Arg	ATC Ile	CCT Pro 390	AAG Lys	GGA Gly	ACG Thr	ACA Thr	CTC Leu 395	ATC Ile	ACC Thr	AAC Asn	CTG Leu	TCA Ser 400	1200
	TCG Ser	GTG Val	CTG Leu	AAG Lys	GAT Asp 405	GAG Glu	GCC Ala	GTC <b>V</b> al	TGG Trp	GAG Glu 410	AAG Lys	CCC Pro	TTC Phe	CGC <b>A</b> rg	TTC Phe 415	CAC His	1248
25	CCC Pro	GAA Glu	CAC His	TTC Phe 420	CTG Leu	GAT Asp	GCC Ala	CAG Gln	GGC Gly 425	CAC His	TTT Phe	GTG Val	AAG Lys	CCG Pro 430	G <b>A</b> G Glu	GCC Ala	1296
30	TTC Phe	CTG Leu	CCT Pro 435	TTC Phe	TCA Ser	GCA Ala	GGC Gly	CGC Arg 440	CGT Arg	GCA Ala	TGC Cys	CTC Leu	GGG Gly 445	GAG Glu	CCC Pro	CTG Leu	1344
	GCC Ala	CGC Arg 450	ATG Met	G <b>A</b> G Glu	CTC Leu	TTC Phe	CTC Leu 455	TTC Phe	TTC Phe	ACC Thr	TCC Ser	CTG Leu 460	CTG Leu	CAG Gln	CAC His	TTC Phe	1392
35	AGC Ser 465	TTC Phe	TCG Ser	GTG Val	CCC Pro	ACT Thr 470	GGA Gly	CAG Gln	CCC Pro	CGG Arg	CCC Pro 475	AGC Ser	CAC His	CAT His	GGT Gly	GTC Val 480	1440
<b>4</b> 0	TTT Phe	GCT Ala	TTC Phe	CTG Leu	GTG Val 485	ACC Thr	CCA Pro	TCC Ser	CCC Pro	TAT Tyr 490	GAG Glu	CTT Leu	TGT Cys	GCT Ala	GTG Val 495	CCC Pro	1488
	CGC <b>A</b> rg	TAG															1494

**4**5 (2) INFORMATION FOR SEQ ID NO: 32:

(i) SEQUENCE CHARACTERISTICS:(A) LENGTH: 497 amino acids(B) TYPE: amino acid

50

#### (D) TOPOLOGY: linear

(ii) 1	MOLECULE	TYPE:	protein
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		(11	) MO	TECO	DE I	IPE:	pro	cem								
5		(xi	) SE	QUEN	CE D	ESCR	IPTI	ON:	SEQ	ID N	0: 3	2 :				
	Met 1	Gly	Leu	Glu	Ala 5	Leu	Val	Pro	Leu	Ala 10	Val	Ile	Val	Ala	Ile 15	Phe
10	Leu	Leu	Leu	Val 20	Asp	Leu	Met	His	Arg 25	Arg	Gln	Arg	Trp	Ala 30	Ala	Arg
	Tyr	Pro	Pro 35	Gly	Pro	Leu	Pro	Leu 40	Pro	Gly	Leu	Gly	Asn 45	Leu	Leu	His
15	Val	Asp 50	Phe	Gln	Asn	Thr	Pro 55	Tyr	Cys	Phe	Asp	Gln 60	Leu	Arg	Arg	Arg
	Phe 65	Gly	Asp	Val	Phe	Ser 70	Leu	Gln	Leu	Ala	Trp 75	Thr	Pro	Val	Val	Val 80
20	Leu	Asn	Gly	Leu	Ala 85	Ala	Val	Arg	Glu	Ala 90	Leu	Val	Thr	His	Gly 95	Glu
	Asp	Thr	Ala	Asp 100	Arg	Pro	Pro	Val	Pro 105	Ile	Thr	Gln	Ile	Leu 110	Gly	Phe
25	Gly	Pro	Arg 115	Ser	Gln	Gly	Val	Phe 120	Leu	Ala	Arg	Tyr	Gly 125	Pro	Ala	Trp
	Arg	Glu 130	Gln	Arg	Arg	Phe	Ser 135	Val	Ser	Thr	Leu	Arg 140	Asn	Leu	Gly	Leu
30	Gly 145	Lys	Lys	Ser	Leu	Glu 150	Gln	Trp	Val	Thr	Glu 155	Glu	Ala	Ala	Суѕ	Leu 160
	Суѕ	Ala	Ala	Phe	Ala 165	Asn	His	Ser	Gly	Arg 170	Pro	Phe	Arg	Pro	Asn 175	Gly
35	Leu	Leu	Asp	Lys 180	Ala	Val	Ser	Asn	Val 185	Ile	Ala	Ser	Leu	Thr 190	Cys	Gly ,
	Arg	Arg	Phe 195	Glu	Tyr	Asp	Asp	Pro 200	Arg	Phe	Leu	Arg	Leu 205	Leu	qaA	Leu
40	Ala	Gln 210	Glu	Gly	Leu	Lys	Glu 215	Glu	Ser	Gly	Phe	Leu 220	Arg	Glu	Val	Leu
	Asn 225	Ala	Val	Pro	Val	Leu 230	Leu	His	Ile	Pro	Ala 235	Leu	Ala	Gly	Lys	Val 240
<b>4</b> 5	Leu	Arg	Phe	Gln	Lys 245	Ala	Phe	Leu	Thr	Gln 250	Leu	Asp	Glu	Leu	Leu 255	Thr
	Glu	His	Arg	Met 260	Thr	Trp	Asp	Pro	Ala 265	Gln	Pro	Pro	Arg	Asp 270	Leu	Thr
50	Glu	Ala	Phe 275	Leu	Ala	Glu	Met	Glu 280	Lys	Ala	Lys	Gly	Asn 285	Pro	Glu	Ser

	Ser	Phe 290	Asn	Asp	Glu	Asn	Leu 295	Cys	Ile	Val	Val	Ala 300	Asp	Leu	Phe	Ser
5	Ala 305	Gly	Met	Val	Thr	Thr 310	Ser	Thr	Thr	Leu	Ala 315	Trp	Gly	Leu	Leu	Leu 320
	Met	Ile	Leu	His	Pro 325	Asp	Val	Gln	Arg	Arg 330	Val	Gln	Gln	Glu	Ile 335	Asp
10	Asp	Val	Ile	Gly 340	Gln	Val	Arg	Arg	Pro 345	Glu	Met	Gly	Asp	Gln 350	Ala	His
15	Met	Pro	Tyr 355	Thr	Thr	Ala	Val	Ile 360	His	Glu	Val	Gln	Arg 365	Phe	Gly	Asp
75	Ile	Val 370	Pro	Leu	Gly	Val	Thr 375	His	Met	Thr	Ser	Arg 380	Asp	Ile	Glu	Val
20	Gln 385	Gly	Phe	Arg	Ile	Pro 390	Lys	Gly	Thr	Thr	Leu 395	Ile	Thr	Asn	Leu	Ser 400
	Ser	Val	Leu	Lys	Asp 405	Glu	Ala	Val	Trp	Glu 410	Lys	Pro	Phe	Arg	Phe 415	His
25	Pro	Glu	His	Phe 420	Leu	Asp	Ala	Gln	Gly 425	His	Phe	Val	Lys	Pro 430	Glu	Ala
	Phe	Leu	Pro 435	Phe	Ser	Ala	Gly	Arg 440	Arg	Ala	Cys	Leu	Gly 445	Glu	Pro	Leu
30	Ala	Arg 450	Met	Glu	Leu	Phe	Leu 455	Phe	Phe	Thr	Ser	Leu 460	Leu	Gln	His	Phe
	Ser 465	Phe	Ser	Val	Pro	Thr 470	Gly	Gln	Pro	Arg	Pro 475	Ser	His	His	Gly	Val 480
35	Phe	Ala	Phe	Leu	Val 485	Thr	Pro	Ser	Pro	Tyr 490	Glu	Leu	Cys	Ala	Val 495	Pro
	Arg															
40	(2)	INFO	ORMAT	rion	FOR	SEQ	ID N	NO: 3	33:							
<b>4</b> 5		(i)	( <i>I</i> (E ()	QUENCA) LE B) TY C) ST C) TO	ENGTH (PE : [RAN]	H: 14 nucl	194 h Leic ESS:	ase acio doul	pai:	cs						
50		(ix)	(1	ATURI A) NA B) L(	AME/I			1491								

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 33:

		( /																
5	ATG Met 1	GGG Gly	CTA Leu	GAA Glu	GCA Ala 5	CTG Leu	GTG Val	CCC Pro	CTG Leu	GCC Ala 10	GTG Val	ATA Ile	GTG Val	GCC Ala	ATC Ile 15	TTC Phe	4	8
	CTG Leu	CTC Leu	CTG Leu	GTG Val 20	GAC Asp	CTG Leu	ATG Met	CAC His	CGG Arg 25	CGC Arg	CAA Gln	CGC Arg	TGG Trp	GCT Ala 30	GCA Ala	CGC Arg	9	6
10	TAC Tyr	CCA Pro	CCA Pro 35	GGC Gly	CCC Pro	CTG Leu	CCA Pro	CTG Leu 40	CCC Pro	GGG Gly	CTG Leu	GGC Gly	AAC Asn 45	CTG Leu	CTG Leu	CAT His	14	4
15	GTG Val	GAC Asp 50	TTC Phe	CAG Gln	AAC Asn	ACA Thr	CCA Pro 55	TAC Tyr	TGC Cys	TTC Phe	GAC Asp	CAG Gln 60	TTG Leu	CGG <b>A</b> rg	CGC <b>A</b> rg	CGC Arg	19	2
	TTC Phe 65	GGG Gly	GAC Asp	GTG <b>V</b> al	TTC Phe	AGC Ser 70	CTG Leu	CAG Gln	CTG Leu	GCC Ala	TGG Trp 75	ACG Thr	CCG Pro	GTG Val	GTC Val	GTG Val 80	24	0
20	CTC eu	AAT Asn	GGG Gly	CTG Leu	GCG Ala 85	GCC Ala	G <b>T</b> G Val	CGC Arg	GAG Glu	GCG Ala 90	CTG Leu	GTG Val	ACC Thr	CAC His	GGC Gly 95	GAG Glu	28	8
25	GAC Asp	ACC Thr	GCC Ala	GAC Asp 100	CGC Arg	CCG Pro	CCT Pro	GTG Val	CCC Pro 105	ATC Ile	ACC Thr	CAG Gln	ATC Ile	CTG Leu 110	GGT Gly	TTC Phe	33	6
	GGG Gly	CCG Pro	CGT Arg 115	TCC Ser	CAA Gln	GGG Gly	GTG Val	TTC Phe 120	CTG Leu	GCG Ala	CGC Arg	TAT Tyr	GGG Gly 125	CCC Pro	GCG Ala	TGG Trp	38	4
30	CGC Arg	GAG Glu 130	CAG Gln	AGG Arg	CGC Arg	TTC Phe	TCC Ser 135	GTC Val	TCC Ser	ACC Thr	TTG Leu	CGC Arg 140	AAC Asn	TTG Leu	GGC Gly	CTG Leu	43	2
25	GGC Gly 145	AAG Lys	<b>AA</b> G	TCG Ser	CTG Leu	GAG Glu 150	CAG Gln	TGG Trp	GTG Val	ACC Thr	GAG Glu 155	G <b>A</b> G Glu	GCC Ala	GCC Ala	TGC Cys	CTT Leu 160	48	0
35	TGT Cys	GCC Ala	GCC <b>A</b> la	TTC Phe	GCC Ala 165	AAC Asn	CAC	TCC Ser	GGA Gly	CGC Arg 170	CCC Pro	TTT Phe	CGC Arg	CCC Pro	AAC Asn 175	GGT Gly	52	8
<b>4</b> 0	CTC Leu	TTG Leu	GAC Asp	AAA Lys 180	GCC Ala	GTG Val	AGC Ser	AAC Asn	GTG Val 185	ATC Ile	GCC Ala	TCC Ser	CTC Leu	ACC Thr 190	TGC Cys	GGG Gly	57	6
	CGC Arg	CGC <b>A</b> rg	TTC Phe 195	GAA Glu	TAC Tyr	GAC Asp	GAC Asp	CCT Pro 200	CGC Arg	TTC Phe	CTC Leu	AGG Arg	CTG Leu 205	CTG Leu	GAC Asp	CTA Leu	62	4
<b>4</b> 5	GCT Ala	CAG Gln 210	GAG Glu	GGA Gly	CTG Leu	AAG Lys	GAG Glu 215	GAG Glu	TCG Ser	GGC Gly	TTT Phe	CTG Leu 220	CGC <b>A</b> rg	GAG Glu	GTG Val	CTG Leu	67	2

55

	AAT Asn 225	GCT Ala	GTC Val	CCC Pro	GTC Val	CTC Leu 230	CTG Leu	CAT His	ATC Ile	CCA Pro	GCG Ala 235	CTG Leu	GCT Ala	GGC Gly	AAG Lys	GTC Val 240	720	)
5	CTA Leu	CGC <b>A</b> rg	TTC Phe	CAA Gln	AAG Lys 245	GCT Ala	TTC Phe	CTG Leu	ACC Thr	CAG Gln 250	CTG Leu	GAT Asp	GAG Glu	CTG Leu	CTA Leu 255	ACT Thr	768	3
40	GAG Glu	CAC His	AGG Arg	ATG Met 260	ACC Thr	TGG Trp	GAC Asp	CCA Pro	GCC Ala 265	C <b>A</b> G Gln	CCC Pro	CCC Pro	CGA Arg	GAC Asp 270	CTG Leu	ACT Thr	816	5
10	GAG Glu	GCC Ala	TTC Phe 275	CTG Leu	GCA Ala	GAG Glu	ATG Met	GAG Glu 280	AAG Lys	GCC <b>A</b> la	AAG Lys	GGG Gly	AAC Asn 285	CCT Pro	GAG Glu	AGC Ser	864	1
15	AGC Ser	TTC Phe 290	AAT Asn	GAT Asp	G <b>A</b> G Glu	AAC Asn	CTG Leu 295	CGC Arg	ATA Ile	GTG Val	GTG Val	GCT Ala 300	GAC Asp	CTG Leu	TTC Phe	TCT Ser	912	2
	GCC Ala :05	GGG Gly	ATG Met	GTG Val	ACC Thr	ACC Thr 310	TCG Ser	ACC Thr	ACG Thr	CTG Leu	GCC Ala 315	TGG Trp	GGC Gly	CTC Leu	CTG Leu	CTC Leu 320	960	)
20	ATG Met	ATC Ile	CTA Leu	CAT His	CCG Pro 325	GAT Asp	GTG Val	CAG Gln	CGC Arg	CGT Arg 330	GTC Val	CAA Gln	CAG Gln	GAG Glu	ATC Ile 335	GAC Asp	1008	3
25	GAC <b>A</b> sp	GTG Val	ATA Ile	GGG Gly 340	CAG Gln	GTG Val	CGG <b>A</b> rg	CGA Arg	CCA Pro 345	GAG Glu	ATG Met	GGT Gly	GAC Asp	CAG Gln 350	GCT Ala	CAC His	1056	5
	ATG Met	CCC Pro	TAC Tyr 355	ACC Thr	ACT Thr	GCC Ala	GTG Val	ATT Ile 360	CAT His	GAG Glu	GTG Val	CAG Gln	CGC Arg 365	TTT Phe	GGG Gly	GAC Asp	1104	1
30	ATC Ile	GTC Val 370	CCC Pro	CTG Leu	GGT Gly	GTG Val	ACC Thr 375	CAT His	ATG Met	ACA Thr	TCC Ser	CGT Arg 380	GAC Asp	ATC Ile	GAA Glu	GTA Val	1157	2
35	CAG Gln 385	GGC Gly	TTC Phe	CGC Arg	ATC Ile	CCT Pro 390	AAG Lys	GGA Gly	ACG Thr	ACA Thr	CTC Leu 395	ATC Ile	ACC Thr	AAC Asn	CTG Leu	TCA Ser 400	1200	D
	TCG Ser	GTG Val	CTG Leu	AAG Lys	GAT Asp 405	GAG Glu	GCC Ala	GTC Val	TGG Trp	GAG Glu 410	AAG Lys	CCC Pro	TTC Phe	CGC <b>Ar</b> g	TTC Phe 415	CAC His	124	8
40	CCC Pro	GAA Glu	CAC His	TTC Phe 420	CTG Leu	GAT Asp	GCC Ala	CAG Gln	GGC Gly 425	CAC His	TTT Phe	GTG Val	AAG Lys	CCG Pro 430	GAG Glu	GCC <b>Al</b> a	129	6
<b>4</b> 5	TTC Phe	CTG Leu	CCT Pro 435	Phe	TCA Ser	GCA Ala	GGC Gly	CGC Arg 440	Arg	GCA Ala	TGC Cys	CTC Leu	GGG Gly 445	Glu	CCC Pro	CTG Leu	134	4
	GCC Ala	CGC Arg 450	Met	GAG Glu	CTC Leu	TTC Phe	CTC Leu 455	Phe	TTC Phe	ACC Thr	TCC Ser	CTG Leu 460	Leu	C <b>A</b> G Gln	CAC His	TTC Phe	139	2

	AGC Ser 465	TTC Phe	TCG Ser	GTG Val	CCC Pro	ACT Thr 470	GGA Gly	CAG Gln	CCC Pro	CGG Arg	CCC Pro 475	AGC Ser	CAC His	CAT His	GGT Gly	GTC Val 480		1440
5	TTT Phe	GCT Ala	TTC Phe	CTG Leu	GTG Val 485	ACC Thr	CCA Pro	TCC Ser	CCC Pro	TAT Tyr 490	GAG Glu	CTT Leu	TGT Cys	GCT Ala	GTG Val 495	CCC Pro		1488
10	CGC Arg	TAG																1494
15	(2)		( E		ENCE ENGTI YPE :	CHAP H: 49	RACTI 97 ar no ac	ERIST mino cid	rics									
20			MOI SEC	LECUI	LE TY	/PE:	prot	cein	SEO 1	ID <b>N</b> (	D: 34	1:						
	Met 1		Leu	-									Val	Ala	Ile 15	Phe		
25	Leu	Leu	Leu	Val 20	Asp	Leu	Met	His	Arg 25	Arg	Gln	Arg	Trp	Ala 30	Ala	Arg		
	Tyr	Pro	Pro 35	Gly	Pro	Leu	Pro	Leu 40	Pro	Gly	Leu	Gly	Asn 45	Leu	Leu	His		
30	Val	Asp 50	Phe	Gln	Asn	Thr	Pro 55	Tyr	Cys	Phe	Asp	Gln 60	Leu	Arg	Arg	Arg		
	Phe 65	Gly	Asp	Val	Phe	Ser 70	Leu	Gln	Leu	Ala	Trp 75	Thr	Pro	Val	Val	Val 80		
35	Leu	Asn	Gly	Leu	Ala 85	Ala	Val	Arg	Glu	Ala 90	Leu	Val	Thr	His	Gly 95	Glu	•	
	Asp	Thr	Ala	Asp 100	Arg	Pro	Pro	Val	Pro 105	Ile	Thr	Gln	Ile	Leu 110	Gly	Phe		
<b>4</b> 0	Gly	Pro	Arg 115	Ser	Gln	Gly	Val	Phe 120	Leu	Ala	Arg	Tyr	Gly 125	Pro	Ala	Trp		
	Arg	Glu 130	Gln	Arg	Arg	Phe	Ser 135	Val	Ser	Thr	Leu	Arg 140	Asn	Leu	Gly	Leu		
<b>4</b> 5	Gly 145	Lys	Lys	Ser	Leu	Glu 150	Gln	Trp	Val	Thr	Glu 155	Glu	Ala	Ala	Cys	Leu 160		
	Cys	Ala	Ala	Phe	Ala 165	Asn	His	Ser	Gly	Arg 170	Pro	Phe	Arg	Pro	Asn 175	Gly		

	Leu	Leu	Asp	Lys 180	Ala	Val	Ser	Asn	Val 185	Ile	Ala	Ser	Leu	Thr 190	Cys	Gly
5	Arg	Arg	Phe 195	Glu	Tyr	Asp	Asp	Pro 200	Arg	Phe	Leu	Arg	Leu 205	Leu	Asp	Leu
	Ala	Gln 210	Glu	Gly	Leu	Lys	Glu 215	Glu	Ser	Gly	Phe	Leu 220	Arg	Glu	Val	Leu
10	Asn 225	Ala	Val	Pro	Val	Leu 230	Leu	His	Ile	Pro	Ala 235	Leu	Ala	Gly	Lys	Val 240
	Leu	Arg	Phe	Gln	Lys 245	Ala	Phe	Leu	Thr	Gln 250	Leu	Asp	Glu	Leu	Leu 255	Thr
15	Glu	His	Arg	Met 260	Thr	Trp	Asp	Pro	Ala 265	Gln	Pro	Pro	Arg	Asp 270	Leu	Thr
	Glu	Ala	Phe 275	Leu	Ala	Glu	Met	Glu 280	Lys	Ala	Lys	Gly	Asn 285	Pro	Glu	Ser
20	Ser	Phe 290	Asn	Asp	Glu	Asn	Leu 295	Arg	Ile	Val	Val	Ala 300	Asp	Leu	Phe	Ser
	Ala 305	Gly	Met	Val	Thr	Thr 310	Ser	Thr	Thr	Leu	Ala 315	Trp	Gly	Leu	Leu	Leu 320
25	Met	Ile	Leu	His	Pro 325	Asp	Val	Gln	Arg	Arg 330	Val	Gln	Gln	Glu	Ile 335	Asp
	Asp	Val	Ile	Gly 340	Gln	Val	Arg	Arg	Pro 345	Glu	Met	Gly	Asp	Gln 350	Ala	His
30	Met	Pro	Tyr 355	Thr	Thr	Ala	Val	Ile 360	His	Glu	Val	Gln	Arg 365	Phe	Gly	Asp
35	Ile	<b>V</b> al 370	Pro	Leu	Gly	Val	Thr 375	His	Met	Thr	Ser	Arg 380	Asp	Ile	Glu	Val
	Gln 385	Gly	Phe	Arg	Ile	Pro 390	Lys	Gly	Thr	Thr	Leu 395	Ile	Thr	Asn	Leu	Ser 400
40	Ser	Val	Leu	Lys	<b>Asp</b> 405	Glu	Ala	Val	Trp	Glu 410	Lys	Pro	Phe	Arg	Phe 415	His
	Pro	Glu	His	Phe 420	Leu	Asp	Ala	Gln	Gly 425	His	Phe	Val	Lys	Pro 430	Glu	Ala
<b>4</b> 5	Phe	Leu	Pro <b>4</b> 35	Phe	Ser	Ala	Gly	Arg 440	Arg	Ala	Cys	Leu	Gly 445	Glu	Pro	Leu
	Ala	Arg 450	Met	Glu	Leu	Phe	Leu 455	Phe	Phe	Thr	Ser	Leu 460	Leu	Gln	His	Phe
50	Ser 465	Phe	Ser	Val	Pro	Thr 470	Gly	Gln	Pro	Arg	Pro 475	Ser	His	His	Gly	Val 480

	Phe	Ala	Phe	Leu	Val 485	Thr	Pro	Ser	Pro	Tyr 490	Glu	Leu	Cys	Ala	Val 495	Pro	
	Arg																
5	(2)	INF	ORMA	TION	FOR	SEQ	ID	NO:	35:								
10		(i)	(, () ()	A) LI B) T C) S'	ENGTI YPE : IRANI	HARA H: 10 nuc: DEDNI	194 l leic ESS:	base acio doul	pai: d	rs							
15		(ix)	()		AME/I	KEY: ION:		1491									
		(xi	) SE	QUEN	CE DI	ESCR:	IPTI	ON:	SEQ :	ID N	D: 35	5:					
20	ATG Met	GGG Gly	CTA Leu	GAA Glu	GCA Ala 5	CTG Leu	GTG Val	CCC Pro	CTG Leu	GCC Ala 10	GTG <b>V</b> al	ATA Ile	GTG Val	GCC Ala	ATC Ile 15	TTC Phe	4.8
	CTG Leu	CTC Leu	CTG Leu	GTG Val 20	GAC Asp	CTG Leu	ATG Met	CAC His	CGG Arg 25	CGC <b>A</b> rg	CAA Gln	CGC Arg	TGG Trp	GCT Ala 30	GCA Ala	CGC Arg	96
25	TAC Tyr	CCA Pro	CCA Pro 35	GGC Gly	CCC Pro	CTG Leu	CCA Pro	CTG Leu 40	CCC Pro	GGG Gly	CTG Leu	GGC Gly	AAC Asn 45	CTG Leu	CTG Leu	CAT His	144
30										TTC Phe							192
	TTC Phe 65	GGG Gly	GAC Asp	GTG Val	TTC Phe	AGC Ser 70	CTG Leu	CAG Gln	CTG Leu	GCC Ala	TGG Trp 75	ACG Thr	CCG Pro	GTG Val	GTC Val	GTG Val 80	240
35										GCG Ala 90							288
40	GAC Asp	ACC Thr	GCC Ala	GAC Asp 100	CGC <b>A</b> rg	CCG Pro	CCT Pro	GTG <b>Val</b>	CCC Pro 105	ATC Ile	ACC Thr	CAG Gln	ATC Ile	CTG Leu 110	GGT Gly	TTC Phe	336
	GGG Gly	CCG Pro	CGT Arg 115	TCC Ser	CAA Gln	GGG Gly	GTG Val	TTC Phe 120	CTG Leu	GCG Ala	CGC Arg	TAT Tyr	GGG Gly 125	CCC Pro	GCG Ala	TGG Trp	384
<b>4</b> 5	CGC Arg	GAG Glu 130	CAG Gln	AGG Arg	CGC Arg	TTC Phe	TCC Ser 135	GTC Val	TCC Ser	ACC Thr	TTG Leu	CGC Arg 140	AAC Asn	TTG Leu	GGC Gly	CTG Leu	432
50																	

											GAG Glu 155					CTT Leu 160	480
5											CCC Pro						528
10											GCC Ala						576
											CTC Leu						624
15											TTT Phe			_			672
20											GCG Ala 235						720
											CTG Leu						768
25											CCC Pro						816
30	GAG Glu	GCC Ala	TTC Phe 275	CTG Leu	GCA Ala	GAG Glu	ATG Met	GAG Glu 280	AAG Lys	GCC Ala	AAG Lys	GGG Gly	AAC Asn 285	CCT Pro	GAG Glu	AGC Ser	864
											GTG Val						912
35											GCC Ala 315						960
40											GTC Val						1008
<b>4</b> 0			Ile		Gln	Val	Arg	Arg	Pro	Glu	ATG Met	Gly	Asp	Gln	Ala		1056
<b>4</b> 5											GTG Val						1104
											TCC Ser						1152

							AAG Lys										1200
5	TCG Ser	GTG Val	CTG Leu	AAG Lys	GAT Asp 405	GAG Glu	GCC Ala	GTC Val	TGG Trp	GAG Glu 410	AAG Lys	CCC Pro	TTC Phe	CGC Arg	TTC Phe 415	CAC His	1248
10							GCC Ala										1296
							GGC Gly										1344
15	GCC Ala	CGC Arg 450	ATG Met	GAG Glu	CTC Leu	TTC Phe	CTC Leu 455	TTC Phe	TTC Phe	ACC Thr	TCC Ser	CTG Leu 460	CTG Leu	CAG Gln	CAC His	TTC Phe	1392
20	AGC Ser 465	TTC Phe	TCG Ser	GTG Val	CCC Pro	ACT Thr 470	GGA Gly	CAG Gln	CCC Pro	CGG Arg	CCC Pro 475	AGC Ser	CAC His	CAT His	GGT Gly	GTC Val 480	1440
							CCA Pro										1488
25	CGC <b>Ar</b> g	TAG															1494

#### (2) INFORMATION FOR SEQ ID NO: 36:

(i) SEQUENCE CHARACTERISTICS:

(A) LENGTH: 497 amino acids

(B) TYPE: amino acid (D) TOPOLOGY: linear

(ii) MOLECULE TYPE: protein

(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 36:

Met Gly Leu Glu Ala Leu Val Pro Leu Ala Val Ile Val Ala Ile Phe

Leu Leu Leu Val Asp Leu Met His Arg Arg Gln Arg Trp Ala Ala Arg 40

Tyr Pro Pro Gly Pro Leu Pro Leu Pro Gly Leu Gly Asn Leu Leu His

Val Asp Phe Gln Asn Thr Pro Tyr Cys Phe Asp Gln Leu Arg Arg Arg 45

Phe Gly Asp Val Phe Ser Leu Gln Leu Ala Trp Thr Pro Val Val Val

50

30

35

	Leu	Asn	Gly	Leu	Ala 85	Ala	Val	Arg	Glu	Ala 90	Leu	Val	Thr	His	Gly 95	Glu
5	Asp	Thr	Ala	Asp 100	Arg	Pro	Pro	Val	Pro 105	Ile	Thr	Gln	Ile	Leu 110	Gly	Phe
	Gly	Pro	Arg 115	Ser	Gln	Gly	Val	Phe 120	Leu	Ala	Arg	Tyr	Gly 125	Pro	Ala	Trp
10	Arg	Glu 130	Gln	Arg	Arg	Phe	Ser 135	Val	Ser	Thr	Leu	Arg 140	Asn	Leu	Gly	Leu
	Gly 145	Lys	Lys	Ser	Leu	Glu 150	Gln	Trp	Val	Thr	Glu 155	Glu	Ala	Ala	Cys	Leu 160
15	Cys	Ala	Ala	Phe	Ala 165	Asn	His	Ser	Gly	Arg 170	Pro	Phe	Arg	Pro	Asn 175	Gly
00	Leu	Leu	Asp	Lys 180	Ala	Val	Ser	Asn	Val 185	Ile	Ala	Ser	Leu	Thr 190	Cys	Gly
20	Arg	Arg	Phe 195	Glu	Tyr	Asp	Asp	Pro 200	Arg	Phe	Leu	Arg	Leu 205	Leu	Asp	Leu
25	Ala	Gln 210	Glu	Gly	Leu	Lys	Glu 215	Glu	Ser	Gly	Phe	Leu 220	Arg	Glu	Val	Leu
	Asn 225	Ala	Val	Pro	Val	Leu 230	Leu	His	Ile	Pro	Ala 235	Leu	Ala	Gly	Lys	Val 240
30	Leu	Arg	Phe	Gln	Lys 245	Ala	Phe	Leu	Thr	Gln 250	Leu	Asp	Glu	Leu	Leu 255	Thr
	Glu	His	Arg	Met 260	Thr	Trp	Asp	Pro	Ala 265	Gln	Pro	Pro	Arg	Asp 270	Leu	Thr
35	Glu	Ala	Phe 275	Leu	Ala	Glu	Met	Glu 280	Lys	Ala	Lys	Gly	Asn 285	Pro	Glu	Ser
	Ser	Phe 290	Asn	Asp	Glu	Asn	<b>Leu</b> 295	Arg	Ile	Val	Val	Ala 300	Asp	Leu	Phe	Ser
40	<b>A</b> la 305	Gly	Met	Val	Thr	Thr 310	Ser	Thr	Thr	Leu	<b>Ala</b> 315	Trp	Gly	Leu	Leu	Leu 320
	Met	Ile	Leu	His	Pro 325	Asp	Val	Gln	Arg	Arg 330	Val	Gln	Gln	Glu	Ile 335	Asp
45	Asp	Val	Ile	Gly 340	Gln	Val	Arg	Arg	Pro 345	Glu	Met	Gly	Asp	Gln 350	Ala	His
	Met	Pro	Tyr 355	Thr	Thr	Ala	Val	Ile 360	His	Glu	Val	Gln	Arg 365	Phe	Gly	Asp
50	Ile	Val 370	Pro	Leu	Gly	Val	Thr 375	His	Met	Thr	Ser	Arg 380	Asp	Ile	Glu	Val

	Gln 385	Gly	Phe	Arg	Ile	Pro 390	Lys	Gly	Thr	Thr	Leu 395	Ile	Thr	Asn	Leu	Ser 400	
5	Ser	Val	Leu	Lys	Asp 405	Glu	Ala	Val	Trp	Glu 410	Lys	Pro	Phe	Arg	Phe 415	His	
	Pro	Glu	His	Phe 420	Leu	Asp	Ala	Gln	Gly 425	His	Phe	Val	Lys	Pro 430	Glu	Ala	
10	Phe	Leu	Pro 435	Phe	Ser	Ala	Gly	Arg 440	Arg	Ala	Cys	Leu	Gly 445	Glu	Pro	Leu	
	Ala	Arg 450	Met	Glu	Leu	Phe	Leu 455	Phe	Phe	Thr	Ser	Leu 460	Leu	Gln	His	Phe	
15	Ser 465	Phe	Ser	Val	Pro	Thr 470	Gly	Gln	Pro	Arg	Pro 475	Ser	His	His	Gly	Val 480	
	Phe	Ala	Phe	Leu	Val 485	Ser	Pro	Ser	Pro	Tyr 490	Glu	Leu	Cys	Ala	Val 495	Pro	
20	Arg																
	(2)			TION													
25		(1)	( <i>I</i> (E	QUENCA) LE B) TY C) ST C) TC	ENGTI (PE : [RANI	H: 14 nucl	194 h Leic	ase acio doul	pain I	cs							
30		(ix)	(7	ATURE A) NA B) LO	ME/I			1491									
		(xi)	SEÇ	QUENC	CE DE	ESCRI	PTIC	ON: 5	SEQ ]	D NO	D: 37	7:					
35	ATG Met 1	GGG Gly	CTA Leu	GAA Glu	GCA Ala 5	CTG Leu	GTG Val	CCC Pro	CTG Leu	GCC Ala 10	GTG Val	ATA Ile	GTG Val	GCC Ala	ATC Ile 15	TTC Phe	4.8
40	CTG Leu	CTC Leu	CTG Leu	GTG Val 20	GAC Asp	CTG Leu	ATG Met	CAC His	CGG Arg 25	CGC Arg	CAA Gln	CGC Arg	TGG Trp	GCT Ala 30	GCA Ala	CGC <b>A</b> rg	96
	TAC Tyr	CCA Pro	CCA Pro 35	Gly GGC	CCC Pro	CTG Leu	CCA Pro	CTG Leu 40	CCC Pro	GGG Gly	CTG Leu	GGC Gly	AAC Asn 45	CTG Leu	CTG Leu	CAT His	144
<b>4</b> 5	GTG Val	GAC Asp 50	TTC Phe	CAG Gln	AAC Asn	ACA Thr	CCA Pro 55	TAC Tyr	TGC Cys	TTC Phe	GAC Asp	CAG Gln 60	TTG Leu	CGG Arg	CGC <b>Ar</b> g	CGC Arg	192
50	TTC Phe 65	GGG Gly	GAC Asp	GTG Val	TTC Phe	AGC Ser 70	CTG Leu	CAG Gln	CTG Leu	GCC Ala	TGG Trp 75	ACG Thr	CCG Pro	GTG Val	GTC Val	GTG Val 80	240

	CTC Leu	AAT Asn	GGG Gly	CTG Leu	GCG Ala 85	GCC Ala	GTG Val	CGC <b>A</b> rg	GAG Glu	GCG Ala 90	CTG Leu	GTG Val	ACC Thr	CAC His	GGC Gly 95	GAG Glu	288
5	GAC Asp	ACC Thr	GCC Ala	GAC Asp 100	CGC Arg	CCG Pro	CCT Pro	GTG Val	CCC Pro 105	ATC Ile	ACC Thr	CAG Gln	ATC Ile	CTG Leu 110	GGT Gly	TTC Phe	336
	GGG Gly	CCG Pro	CGT Arg 115	TCC Ser	CAA Gln	GGG Gly	GTG Val	TTC Phe 120	CTG Leu	GCG Ala	CGC Arg	TAT Tyr	GGG Gly 125	CCC Pro	GCG Ala	TGG Trp	384
10	CGC Arg	GAG Glu 130	C <b>A</b> G Gln	AGG Arg	CGC <b>Ar</b> g	TTC Phe	TCC Ser 135	GTC Val	TCC Ser	ACC Thr	TTG Leu	CGC Arg 140	AAC Asn	TTG Leu	GGC Gly	CTG Leu	432
15	GGC Gly 145	AAG Lys	AAG Lys	TCG Ser	CTG Leu	GAG Glu 150	CAG Gln	TGG Trp	GTG Val	ACC Thr	GAG Glu 155	GAG Glu	GCC Ala	GCC Ala	TGC Cys	CTT Leu 160	480
	TGT Cys	GCC Ala	GCC Ala	TTC Phe	GCC Ala 165	AAC Asn	CAC His	TCC Ser	GGA Gly	CGC Arg 170	CCC Pro	TTT Phe	CGC Arg	CCC Pro	AAC Asn 175	GGT Gly	528
20	CTC Leu	TTG Leu	GAC Asp	AAA Lys 180	GCC <b>A</b> la	GTG Val	AGC Ser	AAC Asn	GTG Val 185	ATC Ile	GCC Ala	TCC Ser	CTC Leu	ACC Thr 190	TGC Cys	GGG Gly	576
<b>2</b> 5	CGC Arg	CGC <b>A</b> rg	TTC Phe 195	GAA Glu	TAC Tyr	GAC Asp	GAC Asp	CCT Pro 200	CGC Arg	TTC Phe	CTC Leu	AGG Arg	CTG Leu 205	CTG Leu	GAC Asp	CTA Leu	624
	GCT Ala	CAG Gln 210	GAG Glu	GGA Gly	CTG Leu	AAG Lys	GAG Glu 215	GAG Glu	TCG Ser	GGC Gly	TTT Phe	CTG Leu 220	CGC <b>A</b> rg	GAG Glu	GTG Val	CTG Leu	672
30	AAT Asn 225	GCT Ala	GTC Val	CCC Pro	GTC Val	CTC Leu 230	CTG Leu	CAT His	ATC Ile	CCA Pro	GCG Ala 235	CTG Leu	GCT Ala	GGC Gly	AAG Lys	GTC Val 240	720
35	CTA Leu	CGC <b>A</b> rg	TTC Phe	CAA Gln	AAG Lys 245	GCT Ala	TTC Phe	CTG Leu	ACC Thr	CAG Gln 250	CTG Leu	GAT Asp	GAG Glu	CTG Leu	CTA Leu 255	ACT Thr	768
	GAG Glu	CAC His	AGG Arg	ATG Met 260	ACC Thr	TGG Trp	GAC Asp	CCA Pro	GCC Ala 265	CAG Gln	CCC Pro	CCC Pro	CGA Arg	GAC Asp 270	CTG Leu	ACT Thr	816
40	GAG Glu	GCC Ala	TTC Phe 275	CTG Leu	GCA Ala	GAG Glu	ATG Met	GAG Glu 280	AAG Lys	GCC Ala	AAG Lys	GGG Gly	AAC Asn 285	CCT Pro	G <b>A</b> G Glu	AGC Ser	864
<b>4</b> 5	AGC Ser	TTC Phe 290	AAT Asn	GAT Asp	GAG Glu	AAC Asn	CTG Leu 295	TGC Cys	ATA Ile	GTG Val	GTG Val	GCT Ala 300	GAC Asp	CTG Leu	TTC Phe	TCT Ser	912

	GCC Ala 305	GGG Gly	ATG Met	GTG Val	ACC Thr	ACC Thr 310	TCG Ser	ACC Thr	ACG Thr	CTG Leu	GCC Ala 315	TGG Trp	GGC Gly	CTC Leu	CTG Leu	CTC Leu 320	960
5	ATG Met	ATC Ile	CTA Leu	CAT His	CCG Pro 325	GAT Asp	GTG Val	CAG Gln	CGC Arg	CGT Arg 330	GTC <b>V</b> al	CAA Gln	CAG Gln	GAG Glu	ATC Ile 335	GAC Asp	1008
10	GAC Asp	GTG Val	ATA Ile	GGG Gly 340	CAG Gln	GTG Val	CGG Arg	CGA Arg	CCA Pro 345	G <b>A</b> G Glu	ATG Met	GGT Gly	GAC Asp	CAG Gln 350	GCT Ala	CAC His	1056
,,	ATG Met	CCC Pro	TAC Tyr 355	ACC Thr	ACT Thr	GCC Ala	GTG Val	ATT Ile 360	CAT His	G <b>A</b> G Glu	GTG Val	CAG Gln	CGC Arg 365	TTT Phe	GGG Gly	GAC Asp	1104
15	ATC Ile	GTC Val 370	CCC Pro	CTG Leu	GGT Gly	GTG Val	ACC Thr 375	CAT His	ATG Met	ACA Thr	TCC Ser	CGT Arg 380	GAC Asp	ATC Ile	G <b>AA</b> Glu	GTA Val	1152
20	CAG Gln 385	GGC Gly	TTC Phe	CGC <b>A</b> rg	ATC Ile	CCT Pro 390	AAG Lys	GGA Gly	ACG Thr	ACA Thr	CTC Leu 395	ATC Ile	ACC Thr	AAC Asn	CTG Leu	TCA Ser 400	1200
. 0	TCG Ser	GTG Val	CTG Leu	AAG Lys	GAT Asp 405	GAG Glu	GCC Ala	GTC Val	TGG Trp	GAG Glu 410	AAG Lys	CCC Pro	TTC Phe	CGC <b>A</b> rg	TTC Phe 415	CAC His	1248
25	CCC Pro	GAA Glu	CAC His	TTC Phe 420	CTG Leu	GAT Asp	GCC Ala	CAG Gln	GGC Gly 425	CAC His	TTT Phe	GTG Val	AAG Lys	CCG Pro 430	GAG Glu	GCC Ala	1296
	TTC Phe	CTG Leu	CCT Pro 435	TTC Phe	TCA Ser	GCA Ala	GGC Gly	CGC Arg 440	CGT <b>A</b> rg	GCA Ala	TGC Cys	CTC Leu	GGG Gly 445	GAG Glu	CCC Pro	CTG Leu	1344
30	GCC Ala	CGC Arg 450	ATG Met	GAG Glu	CTC Leu	TTC Phe	CTC Leu 455	TTC Phe	TTC Phe	ACC Thr	TCC Ser	CTG Leu 460	CTG Leu	CAG Gln	CAC His	TTC Phe	1392
35	AGC Ser 465	TTC Phe	TCG Ser	GTG Val	CCC Pro	ACT Thr 470	GGA Gly	CAG Gln	CCC Pro	CGG <b>Ar</b> g	CCC Pro 475	AGC Ser	CAC His	CAT His	GGT Gly	GTC Val 480	1440
	TTT Phe	GCT Ala	TTC Phe	CTG Leu	GTG Val 485	AGC Ser	CCA Pro	TCC Ser	CCC Pro	TAT Tyr 490	GAG Glu	CTT Leu	TGT Cys	GCT Ala	GTG Val 495	CCC Pro	1488
<b>4</b> 0	CGC Arg	TAG															1494

(2) INFORMATION FOR SEQ ID NO: 38:

(i) SEQUENCE CHARACTERISTICS:(A) LENGTH: 497 amino acids(B) TYPE: amino acid(D) TOPOLOGY: linear

55

(ii)	MOLECULE	TYPE:	protein	
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(xi) S	SEOUENCE	DESCRIPTION:	SEQ	ID	NO:	38:
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5	Met 1	Gly	Leu	Glu	Ala 5	Leu	Val	Pro	Leu	Ala 10	Val	Ile	Val	Ala	Ile 15	Phe
	Leu	Leu	Leu	Val 20	Asp	Leu	Met	His	Arg 25	Arg	Gln	Arg	Trp	Ala 30	Ala	Arg
10	Туг	Pro	Pro 35	Gly	Pro	Leu	Pro	Leu 40	Pro	Gly	Leu	Gly	Asn 45	Leu	Leu	His
	Val	Asp 50	Phe	Gln	Asn	Thr	Pro 55	Tyr	Cys	Phe	Asp	Gln 60	Leu	Arg	Arg	Arg
15	Phe 65	Gly	Asp	Val	Phe	Ser 70	Leu	Gln	Leu	Ala	Trp 75	Thr	Pro	Val	Val	Val 80
20	Leu	Asn	Gly	Leu	Ala 85	Ala	Val	Arg	Glu	Ala 90	Leu	Val	Thr	His	Gly 95	Glu
	qaA	Thr	Ala	Asp 100	Arg	Pro	Pro	Val	Pro 105	Ile	Thr	Gln	Ile	Leu 110	Gly	Phe
25	Gly	Pro	Arg 115	Ser	Gln	Gly	Val	Phe 120	Leu	Ala	Arg	Tyr	Gly 125	Pro	Ala	Trp
	Arg	Glu 130	Gln	Arg	Arg	Phe	Ser 135	Val	Ser	Thr	Leu	Arg 140	Asn	Leu	Gly	Leu
30	Gly 145	Lys	Lys	Ser	Leu	Glu 150	Gln	Trp	Val	Thr	Glu 155	Glu	Ala	Ala	Cys	Leu 160
	Cys	Ala	Ala	Phe	Ala 165	Asn	His	Ser	Gly	Arg 170	Pro	Phe	Arg	Pro	Asn 175	Gly
35	Leu	Leu	Asp	Lys 180	Ala	Val	Ser	Asn	Val 185	Ile	Ala	Ser	Leu	Thr 190	Cys	Gly
	Arg	Arg	Phe 195	Glu	Tyr	Asp	Asp	Pro 200	Arg	Phe	Leu	Arg	Leu 205	Leu	Asp	Leu
40	Ala	Gln 210	Glu	Gly	Leu	Lys	Glu 215	Glu	Ser	Gly	Phe	Leu 220	Arg	Glu	Val	Leu
	<b>Asn 22</b> 5	Ala	Val	Pro	Val	Leu 230	Leu	His	Ile	Pro	Ala 235	Leu	Ala	Gly	Lys	Val 240
<b>4</b> 5	Leu	Arg	Phe	Gln	Lys 245	Ala	Phe	Leu	Thr	Gln 250	Leu	Asp	Glu	Leu	Leu 255	Thr
	Glu	His	Arg	Met 260	Thr	Trp	Asp	Pro	Ala 265	Gln	Pro	Pro	Arg	Asp 270	Leu	Thr
50	Glu	Ala	Phe		Ala	Glu		Glu 280					Asn 285	Pro	Glu	Ser

	Ser	Phe 290	Asn	Asp	Glu	Asn	Leu 295	Cys	Ile	Val	Val	Ala 300	Asp	Leu	Phe	Ser
5	Ala 305	Gly	Met	Val	Thr	Thr 310	Ser	Thr	Thr	Leu	Ala 315	Trp	Gly	Leu	Leu	Leu 320
	Met	Ile	Leu	His	Pro 325	Asp	Val	Gln	Arg	Arg 330	Val	Gln	Gln	Glu	Ile 335	Asp
10	Asp	Val	Ile	Gly 340	Gln	Val	Arg	Arg	Pro 345	Glu	Met	Gly	Asp	Gln 350	Ala	His
	Met	Pro	Tyr 355	Thr	Thr	Ala	Val	Ile 360	His	Glu	Val	Gln	Arg 365	Phe	Gly	Asp
15	Ile	Val 370	Pro	Leu	Gly	Val	Thr 375	His	Met	Thr	Ser	Arg 380	Asp	Ile	Glu	Val
, 0	Gln 385	Gly	Phe	Arg	Ile	Pro 390	Lys	Gly	Thr	Thr	Leu 395	Ile	Thr	Asn	Leu	Ser 400
20	Ser	Val	Leu	Lys	Asp 405	Glu	Ala	Val	Trp	Glu 410	Lys	Pro	Phe	Arg	Phe 415	His
20	Pro	Glu	His	Phe 420	Leu	Asp	Ala	Gln	Gly 425	His	Phe	Val	Lys	Pro 430	Glu	Ala
0.5	Phe	Leu	Pro 435	Phe	Ser	Ala	Gly	Arg 440	Arg	Ala	Cys	Leu	Gly 445	Glu	Pro	Leu
25	Ala	Arg 450	Met	Glu	Leu	Phe	Leu 455	Phe	Phe	Thr	Ser	Leu 460	Leu	Gln	His	Phe
••	Ser 465	Phe	Ser	Val	Pro	Thr 470	Gly	Gln	Pro	Arg	Pro <b>47</b> 5	Ser	His	His	Gly	Val 480
30	Phe	Ala	Phe	Leu	Val 485	Ser	Pro	Ser	Pro	Tyr 490	Glu	Leu	Cys	Ala	Val 495	Pro
	Arg															
35	(2)	INFO	ORMA'	rion	FOR	SEQ	ID 1	<b>1</b> 0: 3	39:							
		(i)	(1	A) LI	ENGT	I: 34	bas	ISTIC	airs							
<b>4</b> 0			(c	c) s:		DEDNE	ESS:	acio sino ear								
46		(xi)	) SE(	QUEN	CE DI	ESCR	[PTI	ON: S	SEQ I	ID NO	D: 39	9:				
<b>4</b> 5	GGA	ACGCI	ATG (	GTGG'	rgcto	GC A	rgga'	FATG!	A AG	ГG						

	(2) INFORMATION FOR SEQ ID NO: 40:	
5	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 56 base pairs</li> <li>(B) TYPE: nucleic acid</li> <li>(C) STRANDEDNESS: single</li> <li>(D) TOPOLOGY: linear</li> </ul>	
10	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 40:  CTCAAAGATC TATGGCCCTG TGTTCACTCT GTATTTTGGC CTCGAGCGCA TGGTGG	56
		30
15	(2) INFORMATION FOR SEQ ID NO: 41:  (i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 28 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: single  (D) TOPOLOGY: linear	
20	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 41:	
	CCACCATGCG CTCGAGGCCA AAATACAG	28
	(2) INFORMATION FOR SEQ ID NO: 42:	
25 30	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 31 base pairs</li> <li>(B) TYPE: nucleic acid</li> <li>(C) STRANDEDNESS: single</li> <li>(D) TOPOLOGY: linear</li> </ul>	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 42:	
	GGGTTCCCGG GAAATAATCA ATGATAGTGG G	31
35	(2) INFORMATION FOR SEQ ID NO: 43:	
<b>4</b> 0	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 32 base pairs</li> <li>(B) TYPE: nucleic acid</li> <li>(C) STRANDEDNESS: single</li> <li>(D) TOPOLOGY: linear</li> </ul>	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 43:	
<b>4</b> 5	GGATTGTAAG CACCCCTGG ATCCAGATAT GC	32
50		

	(2) INFORMATION FOR SEQ ID NO: 44:	
5	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 34 base pairs</li> <li>(B) TYPE: nucleic acid</li> <li>(C) STRANDEDNESS: single</li> <li>(D) TOPOLOGY: linear</li> </ul>	
10	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 44:	
	CCCAGCTCCA AGTAAGTCAG CTGCAGTGAT TACC	34
	(2) INFORMATION FOR SEQ ID NO: 45:	
15	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 42 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: single  (D) TOPOLOGY: linear	
20		
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 45:	
	GGTGGTACCC TTGGGAATGA GGTAGTTTCT GAATTTAACG TC	42
25	(2) INFORMATION FOR SEQ ID NO: 46:	
23	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 33 base pairs  (B) TYPE: nucleic acid	
30	(C) STRANDEDNESS: single (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 46:	
35	AGTCTAGAAT GGATCCTTTT GTGGTCCTTG TGC	33
	(2) INFORMATION FOR SEQ ID NO: 47:	
40	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 30 base pairs</li> <li>(B) TYPE: nucleic acid</li> <li>(C) STRANDEDNESS: single</li> <li>(D) TOPOLOGY: linear</li> </ul>	
<b>4</b> 5	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 47:	
	CCCAGAGCTC TGTCTCCAGA GTGAAAGGAG	30

	(2) INFORMATION FOR SEQ ID NO: 48:	
5	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 30 base pairs</li> <li>(B) TYPE: nucleic acid</li> <li>(C) STRANDEDNESS: single</li> <li>(D) TOPOLOGY: linear</li> </ul>	
10	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 48:	
	ACAGAGCTCT GGGAGAGGAA AACTCCCTCC	30
	(2) INFORMATION FOR SEQ ID NO: 49:	
15	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 54 base pairs</li> <li>(B) TYPE: nucleic acid</li> <li>(C) STRANDEDNESS: single</li> <li>(D) TOPOLOGY: linear</li> </ul>	
20		
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 49:	
	CCATAGATTT TTGAGAGATT GGTTAAGGAT TTGCTGACAT CCTTAATATC TATC	54
25	(2) INFORMATION FOR SEQ ID NO: 50:	
30	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 30 base pairs</li> <li>(B) TYPE: nucleic acid</li> <li>(C) STRANDEDNESS: single</li> <li>(D) TOPOLOGY: linear</li> </ul>	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 50:	
35	GACCCTCGTC ACTTTCTGGA TGAAGGTGGA	30
	(2) INFORMATION FOR SEQ ID NO: 51:	
<b>4</b> 0	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 36 base pairs</li> <li>(B) TYPE: nucleic acid</li> <li>(C) STRANDEDNESS: single</li> <li>(D) TOPOLOGY: linear</li> </ul>	
<b>4</b> 5	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 51:	
	GAAGTAGTTA CTTTTCTTAA AATTTCCACC TTCATC	36
50		
50		

	(2) INFORMATION FOR SEQ ID NO: 52:	
5	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 37 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: single  (D) TOPOLOGY: linear	
10	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 52:  AAAGAATTCC CCAACCCAGA GATGTTTGAC CCTCGTC	31
	(2) INFORMATION FOR SEQ ID NO: 53:	
15	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 59 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: single  (D) TOPOLOGY: linear	
20		
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 53:	
	GGCCAGGCCC TCTCCCACAC AAATCCGTTT TCCTGCTGAG AAAGGCATGA AGTAGTTAC	59
25	(2) INFORMATION FOR SEQ ID NO: 54:	
20	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 44 base pairs</li> <li>(B) TYPE: nucleic acid</li> <li>(C) STRANDEDNESS: single</li> <li>(D) TOPOLOGY: linear</li> </ul>	
30		
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 54:	
	GAGAGGGCCT GGCCCGCATG GAGCTGTTTT TATTCCTGAC CTTC	44
35	(2) INFORMATION FOR SEQ ID NO: 55:	
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 34 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: gingle	
40	(C) STRANDEDNESS: single (D) TOPOLOGY: linear	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 55:	
<b>4</b> 5	CAGGAGTTGT GTCAAGGTCC TTTGGGTCAA TCAG	34
50		

	(2) INFORMATION FOR SEQ ID NO: 56:	
5	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 64 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: single  (D) TOPOLOGY: linear	
10	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 56:	
10	TTGTCAATGG ATTTGCTTCT GTCCCGCCCT TCTATCAGCT GTGCTTCATT CCTGTCTGAG	60
	GATC	64
15	(2) INFORMATION FOR SEQ ID NO: 57:	
	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 55 base pairs</li> <li>(B) TYPE: nucleic acid</li> <li>(C) STRANDEDNESS: single</li> <li>(D) TOPOLOGY: linear</li> </ul>	
20	(b) foroboot. Ishedi	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 57:	
	CAGAAGCAAA TCCATTGACA ACAGGAGTTG TGTCAAGGTC CTTTGGGTCA ATCAG	55
25	(2) INFORMATION FOR SEQ ID NO: 58:	
	(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 60 base pairs  (B) TYPE: nucleic acid	
30	<pre>(C) STRANDEDNESS: single (D) TOPOLOGY: linear</pre>	
	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 58:	
35	CTCAGACAGG AATGAAGCAC AGCTGATAGA AGGGCGGGAC AGAAGCAAAT CCATTGACAA	60
	(2) INFORMATION FOR SEQ ID NO: 59:	
40	<ul> <li>(i) SEQUENCE CHARACTERISTICS:</li> <li>(A) LENGTH: 32 base pairs</li> <li>(B) TYPE: nucleic acid</li> <li>(C) STRANDEDNESS: single</li> <li>(D) TOPOLOGY: linear</li> </ul>	
<b>4</b> 5	(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 59:	
		2.2
	GCAGCCAGAC CATCTGTGCT TCTTCAGACA GG	32
50		
55		

(2) INFORMATION FOR SEQ ID NO: 60:

5		(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 44 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: single  (D) TOPOLOGY: linear
10		(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 60:
	C	ACCATATTA ACTTCCCTCA CTTCTGTGCT ACATGACAAC AAAG 44
	(	2) INFORMATION FOR SEQ ID NO: 61:
15		(i) SEQUENCE CHARACTERISTICS:  (A) LENGTH: 52 base pairs  (B) TYPE: nucleic acid  (C) STRANDEDNESS: single  (D) TOPOLOGY: linear
20		
		(xi) SEQUENCE DESCRIPTION: SEQ ID NO: 61:
	A	ATTCTTTGT TGTCATGTAG CACAGAAGTG AGGGAAGTTA ATATGGTGGT AC 52
25		
25		
	Cla	ims
30	1.	A method for evaluation of the safety of a chemical compound, which comprises the steps of:  (a) reacting a chemical compound with recombinant yeast cells producing human cytochrome P456 molecular species P450 1A2, P450 2C9, P450 2E1 and P450 3A4 together with a yeast NADPH P450 reductase, which may be in the form of a fused enzyme with each of said human cytochrome P450 molecular species, or with the cell free extracts of the yeast cells; and  (b) analyzing the resulting metabolite to determine the safety of the compound.
35	2.	The method according to claim 1, wherein the recombinant yeast cells are yeast cells transformed with plasmids having a gene coding for human cytochrome P450 1A2, P450 2C9, P450 2E1 or P450 3A4 together with a gene coding for yeast NADPH-P450 reductase.
40	3.	The method according to claim 1 or 2, wherein the recombinant yeast cells are yeast cells transformed with plasmids each of which has a fused gene comprising a gene coding for the human cytochrome P450 molecule on the 5'-terminal and a gene coding for the yeast NADPH-P450 reductase on 3' terminal.
<b>4</b> 5	4.	The method according to any one of claims 1 to 3, wherein the analyzing of the metabolite is carried out by the Ames Test.
50	5.	The method according to claim 4, wherein the test is carried out using His <sup>-</sup> Salmonella or Trp <sup>-</sup> Escherichia coli.
50	6.	The method according to any one of claims 1 to 5, wherein the recombinant yeast cells further product at least one additional human cytochrome P450 molecular species selected from a group of human cytochrome P450 2A6, P450 2C19 and P450 2D6.
55	7.	The method according to any one of claims 1 to 6, wherein the recombinant yeast cells further product at least one additional human cytochrome P450 molecular species selected from a group of human cytochrome P450 1A1, P450 2B6, P450 2C8 and P450 2C18.

- An artificial fused enzyme, which comprises human cytochrome p450 3A4 and yeast NADPH-P450 reductase.
- 9. A yeast expression plasmid having a fused gene comprising a gene coding for human P450 3A4 and a gene coding for the yeast NADPH-P450 reductase.
- 10. A method of determining in vitro the human metabolite of a chemical compound, which comprises the steps of:

(a) reacting a chemical compound with recombinant yeast cells producing human cytochrome P450 molecular species P450 1A2, P450 2C9, P450 2E1 and P450 3A4 together with a yeast NADPH-P450 reductase, which may be in the form of a fused enzyme with each of said human cytochrome P450 molecular species, or with the cell free extracts of the yeast cells; and

(b) identifying the resulting metabolite.

0

1 4 2	5'-CACAGAGCTCCTCCTGGCCTCTGCCATCTTC-3' 5'-TTACAGGCCCTGCACTTGGCTAAAGCTGC-3'	Frimer for amplifying P4501A2 1.5Kb fragment
503	S'-AGTCTAGAATGGATTCTATTGTGTCCCTTGTGCTC-3' S'-CTCCAAACAAGTCAACTGCAGTGTTTTCCAAGC-3'	Primer for amplifying P450209 0.9Kb fragment
	5'-GCTTGGAAAACACTGCAGTTGACTTGTTTGGAG-3' 5'-actgagcagccaggccatctgctttc-3'	Primer for amplifying P450209 0.8Kb fragment
2 2 1	5'-CCCCAGAATTCAATGTCTGCCCTCGGAGTG-3' S'-CCTCTGGATCCGGCTCTCATTGCCCTGTTTC-3'	Primer for amplifying P4502I1 0.5Ib fragment
	S'-GAAACAGGGCAATGAGAGCCGGATCCAGAGG-3' S'-Gaaaacttgtttgcatgcgggggttcagg-3'	Primer for amplifying P450261 1.00b fragment

Fig. 1

3.4.4	5'-AGTAAGGAATCTAGAAATGGCTCTCATCCCAG-3' 5'-accotorigategringagette-1'	Primer for P4503A4 0.8K	Primer for amplifying P4503A4 0.81b fragment
		7	7.7.7.
	5' - CAAAGCTCTGTCCGATCTGGAGCTCGT-3' 5' - CAAAGTAATTTGAGGTACCTGGTGTTCTCAGGC-3'	P4503A4 0.9X	Frimer ior ampiriying P4503A4 0.9Kb fragment
141	5'-CCTCTAGAAATGCTTTTCCCAATCTCCATG-3'	Primer for	Primer for amplifying
	5'-CCAATCACTGTGTCGAGCTCCTTTGGATC-3'	P4501A1 1.0	P4501A1 1.0Kb fragment
	5'-GATCCAAGAGGAGCTCGACACAGTGATTGG-3'	Primer for	Primer for amplifying
	5'-GGGCTCTCAAGCACCTAAGAGGCGCAGCTGC-3'	P4501A1 0.5	P4501A1 0.5Kb fragment
2 A 6	5'-GCTTCTAGAATGCTGGCCTCAGGGATGCTTC-3'	Primer for	Primer for amplifying
	5'-CGTGGAGGTTGACGTGAACTGGAAGATTC-3'	P4502A6 0.6	P4502A6 0.6Kb fragment
	S'-GAATCTTCCAGTTCACGTCAACCTCCACG-3'	Primer for	Primer for amplifying
	S'-AGACCTGGTACCGCACAGCCTCGCTCAG-3'	P4502A6 0.9	P4502A6 0.9Kb fragment

Fig. 2

286	5'-CCTCTAGAAAATGGAACTCAGCGTCCTCCT-3' 5'-GGGGATCCTGAATGACCCTGGAATCCTTTG-3'	Frimer for amplifying P450286 1.5Kb fragment
208	5'-GAAGAGAAGTCTAGAATGGAACCTTTTGTGGTCC-3' 5'-ATAGCAGATCGGCAGCCAGATGGGCTAGCATTC-3'	Primer for amplifying P4502C8 1.5Xb fragment
2 C 1 8	2C18 5'-AGICTAGAAIGGIACCAGCIGIGGCICIGG-3' 5'-CCCCAAACAIAICAGTIACAGIGGCIAICAAGC-3'	Primer for amplifying P4502C18 0.9Kb fragment
	5' -CCCGATTATTGGAAATATCCTGCAGTTAGATG-3' 5' -Acagcacaggcagcaaactatctgcc-3'	Primer for amplifying P4502C18 1.4Kb fragment

Fig. 3

The sequence shown by 5'-..-3' is described In SEQ ID Nos: 20 to 40. 2C19

Primer for amplifying P450206 0.40b frayment Primer for amplifying P450206 0.91b fragment S'-TGTTCAGCCTGCAGCTGGCCTGGAC-3' S'-AAGCGAGGGTGGTCGTATTCGAAGCG-3' 208

S'-GCTTCGAATACGACGACCCTCGCTTCCTC-3' 5'-ACTAGGTACCCCATTCTAGCGGGGGACAG-3'

Fig. 4

Primer for amplifying P4503A4 Xb1-Xh01 fragment

3A4 (An artificial fused enzyme)

<del>ب</del> 5' - AATCTAGAAATGGCTCTCATCCCAG - 3' 5' - AGGACTCGAGGGGCTCCACTTACGGTGCCATCCC - 3

## EP 0 644 267 A2

(1) Linker for cloning 1A2

5' - AGCITAAAAAATGGCATTGTCCCAGTCTGTTCCCTTCTCGGCCACAGAGCT-3'
3' - ATTITITACCGTAACAGGGTCAGACAAGGGAAGAGCGGTGTC -5'

(2) Linker for cloning 2D6

Fig. 5

5'-CTAGATATGGGGCTAGAAGCACTGGTGCCCCTGGCCGTGATAGTGG-3'
3'- TATACCCGGATCTTGGTGACCACGGGACCGGCACTATCACC-5'

5' - CCAICTICCIGCICCIGGIGGACCIGAIGCACCGGCGCCAACGCIGGGCIGCACGCIACCCACCAGGCCCCCTGCCACIGCCACIGCCGGGCIGCA-3'  5' -GGGCTGGGCAACCTGCAGCTGCAGTGGAACTTCCAGAACACACATACTGCTTCGACCAGTTGCGGCGCGCTTCGGGGAACGTTTAGGCTGCTA-3' 3' -CCCGAACCCGTTGGACGAAGCCGTGAAAGCTGTGAAGCCGAAGCCCTGCAAGTCGG -5'

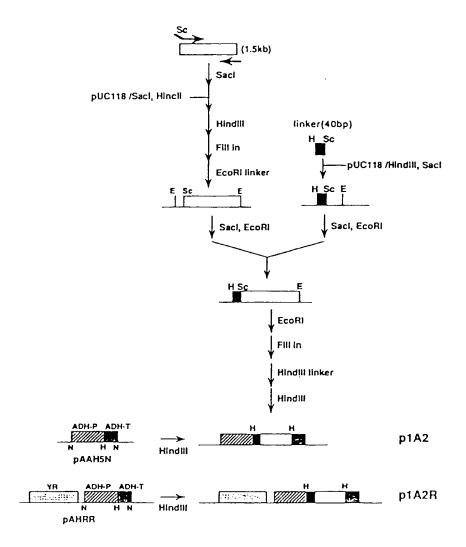


Fig. 6

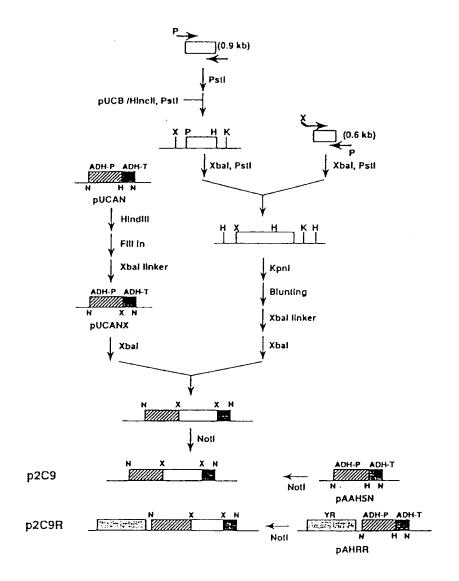


Fig. 7

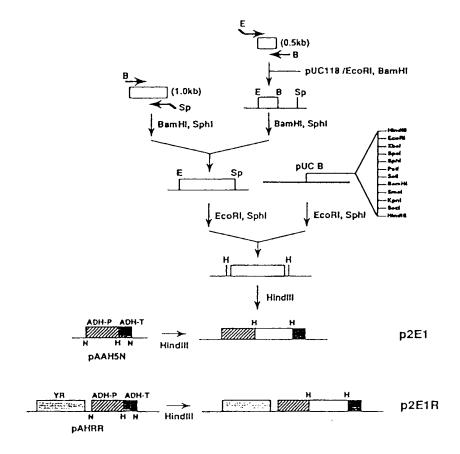


Fig. 8

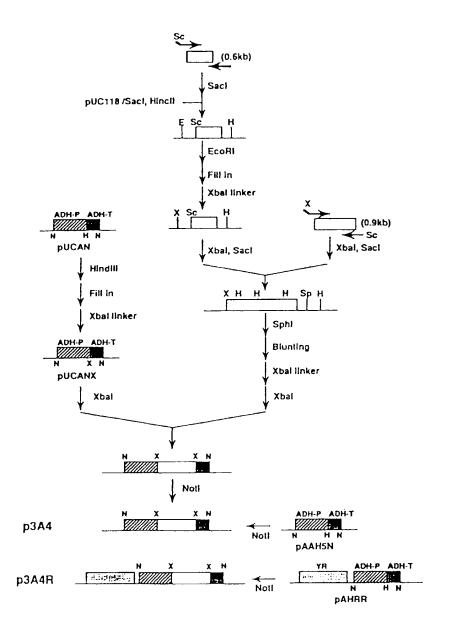


Fig. 9

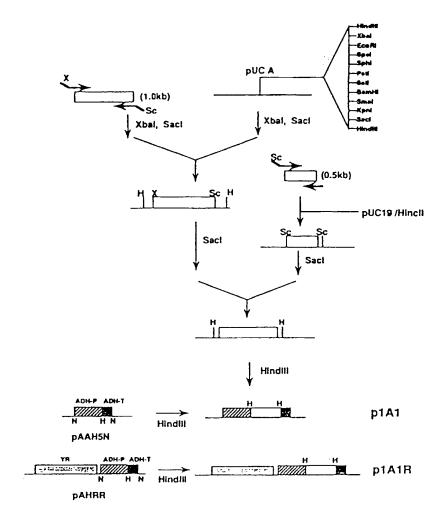


Fig. 10

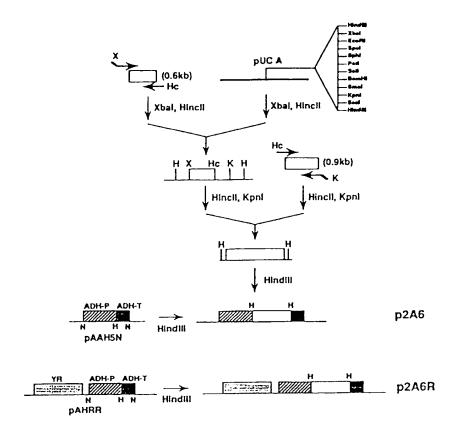


Fig. 11

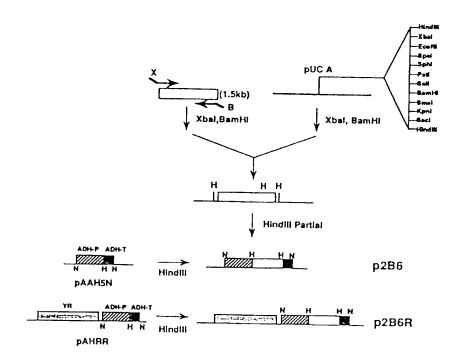


Fig. 12

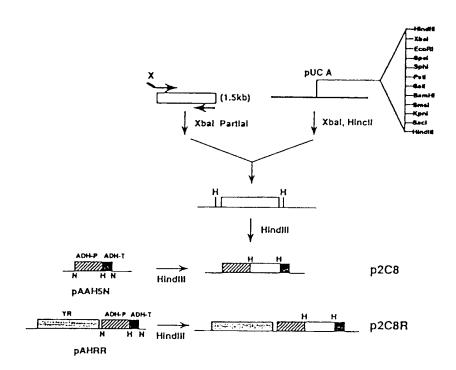


Fig. 13

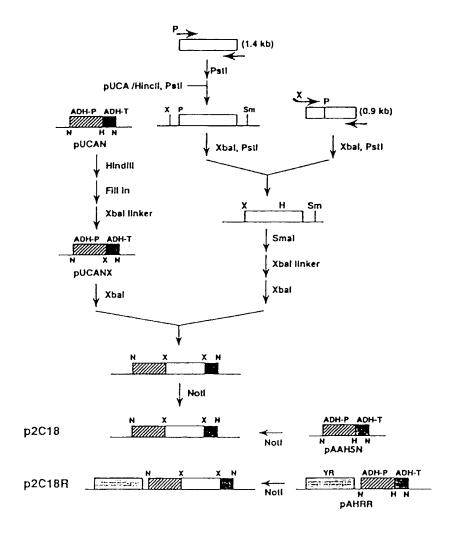


Fig. 14

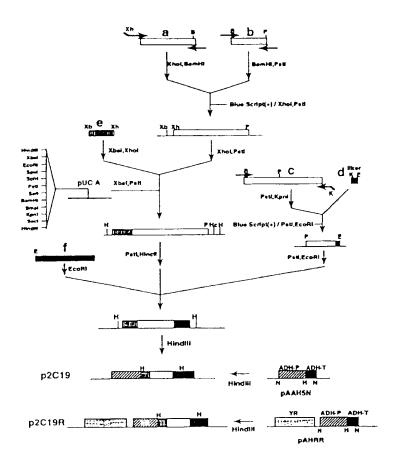


Fig. 15

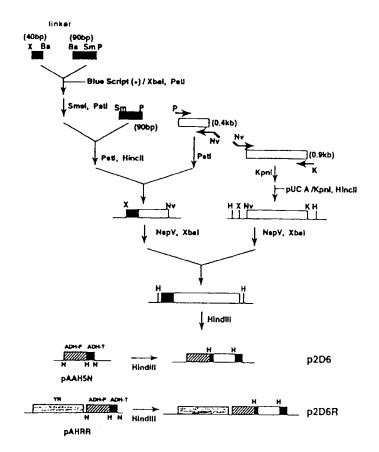


Fig. 16

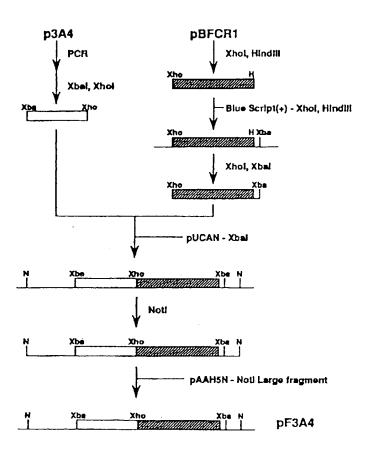


Fig. 17

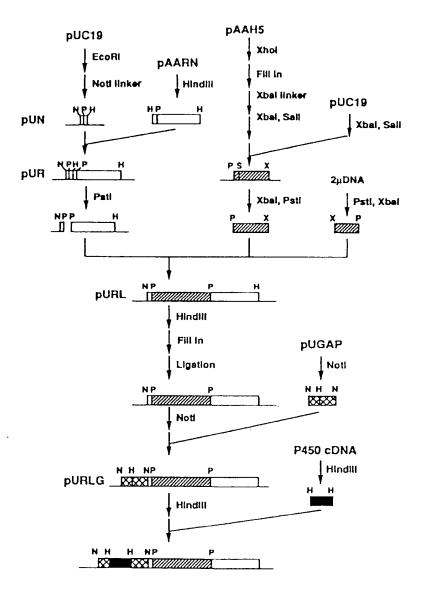


Fig. 18